THE UNITED
REPUBLIC OF
TANZANIA
PRIME MINISTER'S
OFFICE REGIONAL
ADMINISTRATION AND
LOCAL GOVERNMENT
MBEYA REGIONAL
SECRETARIAT

INVITATION FOR TENDERS

TENDER NO.: 78/2025/2026/G/72

FOR

Procurement of medicines from prime vendor for Councils health facilities

19/06/2026

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LIST OF ABBREVIATIONS

AG	Attorney General
Cap	Chapter
CFR	Cost and Freight
DDP	Delivered Duty Paid
EFD	Electronic Financial Devise
ES-MSIP	Environmental Social Management Strategies and Implementation Plan
FY	Financial Year
GCC	General Conditions of Contract
GCLA	Government Chemist Laboratory Authority
GPSA	Government Procurement Services Agency
IFT	Invitation for Tenders
ITT	Instruction to Tenderers
JV	Joint Venture
JVCA	Joint Venture, Consortium, or Association
NCT	National Competitive Tendering
NeST	National Electronics Procurement System of Tanzania
PE	Procuring Entity
PPAA	Public Procurement Appeals Authority
PPRA	Public Procurement Regulatory Authority
SCC	Special Conditions of Contract
SPP	Sustainable Public Procurement
STD	Standard Tender Document
TAEC	Tanzania Atomic Energy Commission
TBS	Tanzania Bureau of Standards
TDS	Tender Data Sheet
TMDA	Tanzania Medicines and Medical Devices Authority

WHO	World Health Organization

PART 1 – TENDERING PROCEDURES

SECTION I: INVITATION NOTICE OF TENDERS

THE UNITED REPUBLIC OF TANZANIA
PRIME MINISTER'S OFFICE REGIONAL ADMINISTRATION AND LOCAL GOVERNMENT
MBEYA REGIONAL SECRETARIAT

Tender No. 78/2025/2026/G/72

FOR

Procurement of medicines from prime vendor for Councils health facilities

19/06/2026

1. This Invitation for Tenders follows the General Procurement Notice for the procurement of Health Sector Goods which appeared in the National e-Procurement System of Tanzania (NeST) dated 18/08/2025.
2.

The Government of Tanzania has set aside funds for the operation of the MBEYA REGIONAL SECRETARIAT during the financial year 2025/2026. It is intended that part of the proceeds of the fund will be used to cover eligible payments under the contract for the Procurement of medicines from prime vendor for Councils health facilities.

3. The MBEYA REGIONAL SECRETARIAT now invites tenders from eligible GOVERNMENT_ENTERPRISE, Company Local, Company Foreign, Special Group, Manufacturer Local, Sole Proprietor Local, Partnership Local,

Manufacturer Foreign, Partnership Foreign and Sole Proprietor Foreign Suppliers of Procurement of medicines from prime vendor for Councils health facilities.

4. Tendering will be conducted through the National Competitive Tendering method specified in the Public Procurement Regulations and is open to specified tenderers in paragraph 3 of this IFT.
5. Interested eligible Tenderers may obtain further information from and inspect the tendering document through NeST. A complete set of tendering document(s) in English may be accessed through NeST.
6. Tenderers are required to register on the NeST and pay tender participation fee indicated in the NeST to be able to participate in this tendering process.
7. All tenders must be accompanied by a Tender Securing Declaration in the format provided in the Tendering Documents.
8. All tenders must be properly filled in and submitted through NeST at or before 10:00 AM hours local time on 21/07/2026. Tenders will be opened promptly thereafter through NeST. Tender opening details will be available to the public through NeST.
9. Tenders not received through NeST shall not be evaluated irrespective of the circumstances.

REGIONAL ADMINISTRATIVE SECRETARY

P.O BOX 754, MBEYA

SECTION II: INVITATION TO TENDERERS (ITT)

A. Introduction

ITT Clause N	Required data/Information	ITT Sub Clause	ITT Clause Description
1.	Scope of Tender and Tendering Method	1.1	The Procuring Entity (PE), as specified in the Tender Data Sheet (**TDS**) and in the Special Conditions of Contract (**SCC**), invites tenders for the supply of Goods Specified in the **TDS** which may include pharmaceuticals, medical equipment and supplies, Laboratory Equipment and reagents, Diagnostic and medical imaging, or nutritional supplements as specified in Section VII - Schedule of Requirements. The successful Tenderer will be expected to supply the goods within the period stated in the **TDS** from the start date specified in the **TDS**. The Contract duration shall be as specified in the **TDS**.
		1.2	Tendering will be conducted through the method of procurement indicated in **TDS** and is open to all Tenderers who meet the eligibility criteria stated in Instructions to Tender (ITT) 3 [Eligible Tenderers]
		1.3	Unless otherwise stated, throughout this tendering document definitions and interpretations shall be as prescribed in the General Conditions of Contract (**GCC).**
2.	Source of Funds	2.1	The Government of Tanzania has set aside funds for the operations of the PE named in the **TDS** during the Financial Year indicated in the **TDS**. It is intended that part of the proceeds of the funds will be applied to cover eligible payments under the contract for the supply of goods as described in the **TDS**. or The Government of Tanzania through the PE named in

			the **TDS** has received/has applied for/intends to apply for a [loan/credit /grant] from the financing institution named in the **TDS** towards the cost of the project, and it intends to apply part of the proceeds of this [loan/credit] to payments under the contract described in the **TDS**.
		2.2	Payments will be made directly by the PE (or by the financing institution specified in the **TDS** upon request by the PE to so pay) for each order and will be subject in all respects to the terms and conditions of the resulting contract placed by the PE.
3.	Eligible Tenderers	3.1	The Invitation for tenders **(IFT)** is open to all tenderers except where it is otherwise specified in the **TDS.** A Tenderer may be natural persons, company(ies) or firms, or public or semi-public agencies of Tanzania and foreign countries, subject to **ITT 3.5** or any combination of them with a formal intent or letter of intent to enter into an agreement or under an existing agreement in the form of a joint venture, consortium, or association (hereinafter referred as JVCA).
		3.2	For a JVCA to be eligible, each member of JVCA is required to be eligible to participate in public procurement and where one party is deemed to be ineligible, the whole JVCA shall be declared ineligible.
		3.3	In the case of a JVCA, all members shall be jointly and severally liable for the execution of the Contract in accordance with the contract terms. The JVCA shall nominate a Lead Member as specified in the **TDS**, who shall have the authority to conduct all business for and on behalf of any and all the members of the JVCA during the tendering process and, in the event the JVCA is awarded the Contract, during contract

			execution. Unless specified in the **TDS**, there is no limit on the number of members in a JVCA.
		3.4	The appointment of a Lead Member in the JVCA shall be confirmed by submission of a valid Power of Attorney to the PE.
		3.5	Any signed agreement that forms a JVCA or a Letter of Intent to execute a JVCA Agreement shall be attested and submitted as part of the tender. Where a Letter of Intent to execute a JVCA is submitted, a draft JVCA Agreement shall also be submitted as part of the tender.
		3.6	Any Tender from a JVCA shall indicate the part of proposed contract to be performed by each party and each party shall be evaluated or post qualified with respect to its contribution only and the responsibilities of each party and shall not be substantially altered without prior written approval of the PE.
		3.7	National Tenderers shall satisfy all relevant licensing and/or registration requirements with the appropriate statutory bodies in Tanzania. Foreign Tenderers are exempted from this requirement but where selected as having submitted the lowest evaluated Tender the successful Tenderer shall register with the appropriate statutory body and shall be required to submit evidence of registration as an approved Supplier in Tanzania before signing the contract.
		3.8	A Tender shall not have a conflict of interest. All Tenderers found to be in conflict of interest shall be disqualified. A Tenderer may be considered to have a conflict of interest with one or more parties in this Tendering process, if they: a) Are associated or have been associated in the past, directly or indirectly with a firm or any of it's affiliates

			which have been engaged by the PE to provide consulting services for the preparation of the specifications and other documents to be used for the procurement of the goods to be procured under this Invitation for Tenders. b) have controlling shareholders in common; or c) receive or have received any direct or indirect subsidy from any of them; or d) have the same legal representative for purposes of this Tender; or e) have a relationship with each other, directly or through common third parties, that puts them in a position to have access to information about or influence on the Tender of another Tenderer, or influence the decisions of the PE regarding this Tendering process; or f) submit more than one Tender in this Tendering process. However, this does not limit the participation of subcontractors in more than one Tender, or as Tenderers and subcontractors simultaneously; or g) participated as a consultant in the preparation of the technical specifications of the goods and services that are the subject of the Tender.
		3.9	A Tenderer may be ineligible if – (a) the Tenderer is declared bankrupt or, in the case of company or firm, insolvent; (b) payments in favour of the Tenderer is suspended in accordance with the judgment of a court of law other than a judgment declaring bankruptcy and resulting, in accordance with the national laws, in the total or partial loss of the right to administer and dispose of its property; (c) legal proceedings are instituted against such Tenderer involving an order suspending payments and

			which may result, in accordance with the national laws, in a declaration of bankruptcy or in any other situation entailing the total or partial loss of the right to administer and dispose of the property; (d) the Tenderer is convicted, by a final judgment, of any offence involving professional conduct; (e) the Tenderer is debarred and blacklisted in accordance with Public Procurement Act, ineligible from participating in public procurement for corrupt, coercive, collusive, fraudulent or obstructive practices, failure to abide by a Tender Securing Declaration, breach of a procurement contract, making false representations about their qualifications during the tender proceedings, or other grounds as may be deemed necessary by the Authority; and f) the tenderer is from an ineligible country as specified in **Section VI** [Eligible Countries] of the tendering document.
		3.10	Public or semi-public owned enterprises in the Government of Tanzania may participate only if they are legally and financially autonomous, if they operate under commercial law by the relevant registration board or authorities and if they are not a dependent agency of the Government.
		3.11	Tenderers shall provide to the PE evidence of their eligibility, proof of compliance with the necessary legal, technical and financial requirements including batch capacity and lead time and their capability and, adequacy of resources to carry out the contract effectively.
		3.12	Tenderers shall provide such evidence of their continued eligibility satisfactory to the PE, as the PE shall reasonably request.

		3.13	Tenderers shall submit proposals relating to the nature, conditions and modalities of sub-contracting wherever the sub-contracting of any elements of the contract amounting to more than ten (10%) percent of the tender price is envisaged.
		3.14	Subject to ITT 3.1 to 3.13, Tenderers must meet the qualification criteria specified under **Section IV** [Qualification and Evaluation Criteria].
4.	Eligible Goods and Related Services	4.1	All goods and related services to be supplied under the contract shall have their origin in eligible source countries, and all expenditures made under the contract will be limited to such goods and related services.
		4.2	For the purposes of this Clause, the term "goods" includes health commodities, raw materials, machinery, equipment and biomedical plants, and "related services" includes services such as insurance, installation, training and initial maintenance.
		4.3	For purposes of this Clause, "origin" means the place where the goods are mined, grown, cultivated, produced, manufactured, or processed, or through manufacture, procession, or assembly, another commercially recognized article results that differs substantially in its basic characteristics from its imported components or the place from which the related services are supplied.
		4.4	The nationality of the firm that produces, assembles, distributes, or sells the goods and services shall not determine their origin.
		4.5	To establish the eligibility of the supplies and the related services, Tenderers shall fill the country of origin declarations included in the Form of Tenders.

		4.6	If so required in the **TDS**, the Tenderer shall demonstrate that it has been duly authorized by the manufacturer of the goods to supply in the United Republic of Tanzania, the goods indicated in its Tender.
5.	One Tender per Tenderer	5.1	A Tenderer shall submit only one tender, in the same Tendering process, either individually or as a partner in a JVCA.
		5.2	No tenderer can be a subcontractor while submitting a tender individually or as a partner of a joint venture in the same Tendering process.
		5.3	A tenderer, if acting in the capacity of subcontractor in any Tender, may participate in more than one Tender but only in that capacity.
		5.4	A Tenderer who submits or participates in more than one tender (other than as a subcontractor or in cases of alternatives that have been permitted or requested) will cause all the tenders in which the Tenderer has participated to be disqualified.
6.	Cost of Tendering	6.1	The Tenderer shall bear all costs associated with the preparation and submission of its tender, and the PE shall in no case be responsible or liable for those costs, except where the PE is ordered by the Public Procurement Appeals Authority (PPAA) to compensate the Tenderer following a successful Tenderer's appeal of the procurement proceedings.

B. Tendering Documents

7.	Content of Tendering Documents	7.1	The goods required, Tendering procedures, and contract terms are prescribed in the Tendering Documents. In addition to the Invitation for Tenders, the Tendering documents, which should be read in conjunction with any addenda issued in accordance with ITT 9.2 [Amendment of Tendering Documents] include: **PART 1: TENDERING PROCEDURES** Section II - Instructions to Tenderers (ITT). Section III - Tender Data Sheet (TDS). Section IV– Qualification and Evaluation Criteria. Section V - Tendering Forms. Section VI - Eligible Countries. **PART 2: PROCURING ENTITY'S REQUIREMENTS** Section VII – Schedule of Requirements **PART 3: CONDITIONS OF CONTRACT AND CONTRACT FORMS** Section VIII - General Conditions of Contract (GCC). Section IX - Special Conditions of Contract (SCC). Section X - Contract Forms.
		7.2	The Invitation for Tenders (IFT) (Section I) issued by the PE is not part of the Tendering Documents and is included as a reference only. In case of discrepancies between the Invitation for Tender and the Tendering Documents listed in ITT 7.1 above, the said Tendering Documents will take precedence.
		7.3	The PE shall not be responsible for any Tendering

			Documents and their addenda, if they were not obtained directly from NeST.
		7.4	The Tenderer is expected to examine all instructions, forms, terms, and specifications in the Tendering documents. Failure to furnish all information required by the Tendering Documents or to submit a tender substantially responsive to the Tendering documents in every respect will be at the Tenderer's risk and may result in the rejection of its tender.
8.	Clarification of Tendering Documents	8.1	A prospective Tenderer requiring any clarification of the Tendering Documents may notify the PE through NeST at least three (3) days for open competitive methods and two (2) days in the case of other tendering methods prior to tender submission deadline.
		8.2	The PE shall within two (2) working days after receiving the request for clarification respond and publish through NeST.
		8.3	PE's response shall include a description of the inquiry without identifying its source.
		8.4	Should the PE deem it necessary to amend the Tendering documents as a result of a clarification, it shall do so following the procedure under **ITT 9** [Amendment of Tendering Documents].
9.	Amendment of Tendering Documents	9.1	Before the deadline for submission of tenders, the PE, for any reason, whether at its own initiative or in response to clarification requested by a prospective Tenderer, may modify the Tendering Documents by issuing Addenda.
		9.2	Any addendum issued, including the notice of any extension of the deadline, shall be part of the Tendering documents pursuant to **ITT 7.1** [Content

			of Tendering Documents] and shall be communicated through NeST to the participating Tenderers
		9.3	In order to allow prospective Tenderers reasonable time to take into account an addendum in preparing their tenders, the PE at its discretion may extend the deadline for submission of tenders, pursuant to **ITT 23.2** [Deadline for Submission of Tenders].

C. Preparation of Tenders

10.	Language of Tender	10.1	The tender prepared by Tenderer, as well as all correspondence and documents relating to the tender exchanged by the Tenderer and the PE, shall be written in the language specified in the **TDS**. Supporting documents and printed literature furnished by the Tenderer may be in another language provided they are accompanied by an accurate translation of the relevant passages in the Language of the Tender, in which case, for purposes of interpretation of the Tender, the translation shall govern.
11.	Documents and Samples Constituting the Tender	11.1	The Tender prepared by the Tenderer shall shall comprise Tendering Forms as listed in **Section V** as follows: a) Form of Tender and a Price Schedule completed in accordance with **ITT 15 [Form of Tender]**, ITT 17 **[Tender Prices and Discounts]**, and ITT 18 **[Tender Currencies]**; b) Sample(s) as requested in the **TDS**; c) Alternative offers if were invited in accordance with Instructions to Tenderers **ITT 16 [AlternativeTenders by Tenderers]**; d) Documentary evidence established in accordance with **ITT13 [Documents Establishing Eligibility and Qualification of the Tenderer]** that the Tenderer is eligible to Tender and is qualified to perform the contract if its Tender is accepted; e) Documentary evidence established in accordance with **ITT 13.3(a)** that the Tender has been authorized by the manufacturer to supply the goods into the United Republic of Tanzania, where required and where the supplier is not the

			manufacturer of those goods. In the case of a Tenderer offering to supply Goods under the Contract that the Tenderer manufactures or otherwise produces (using ingredients supplied by primary manufacturers) that the Tenderer is incorporated in the country of manufacture of the Good, then: (i) has been licensed by the regulatory authority in the country of manufacture to supply the Goods. (ii) has manufactured and marketed the specific Goods covered by this tendering document, for at least two (2) years, and for similar Goods for at least five (5) years. (iii) has received a satisfactory GMP inspection certificate in line with the WHO certification scheme on pharmaceuticals moving in International Commerce from the regulatory authority (RA) in the country of manufacture of the goods or has been certified by the competent authority of a member country of the Pharmaceuticals Inspection Convention (PIC) and has demonstrated compliance with the quality standards during the past two years prior to Tender submission.(ii) that, in the case of a Tenderer offering to supply Goods under the Contract that the Tenderer does not manufacture or otherwise produce: Likewise, if tenderer (ii) has been duly authorized by a manufacturer of the Goods that meets the criteria under to supply the Goods in the Purchaser's Country; shall also submit the following additional information: (a) copies of its audited financial statements for the past three fiscal years. (b) list of major supply contracts conducted within the last five years; f) Documentary evidence established in

			accordance with **ITT12 [Documents Establishing Eligibility of Goods and Related Services and Conformity to Tendering Documents]** that the goods and ancillary services to be supplied by the Tenderer are eligible goods and services and conform to the Tendering Documents; g) Tender Security or Tender Securing Declaration furnished in accordance with ITT19 [Tender Security or Tender Securing Declaration]; h) Code of Conduct for Supplier's Personnel (Environmental and Social - ES) as defined in Sub- Clause 1 (ii) of the General Conditions of Contract, to ensure compliance with the Contractor's ES obligations under the Contract. The Tenderer shall use for this purpose the Code of Conduct form provided in **Section V [Tendering Forms]**. No substantial modifications shall be made to this form, except that the Tenderer may introduce additional requirements, including as necessary to take into account specific contract issues/risks. i) Management Strategies and Implementation Plans (MSIPs) to manage ES risks in a form provided in **Section V [Tendering Forms]**. j) Duly Notarized Power of Attorney (in the format provided in **Section V [Tendering Forms]** authorizing the signatory of the Tender to commit the Tenderer or an Extract of Business Name Registration in case of a Sole Proprietor in accordance with **ITT 20.2 [Format and Signing of Tenders]** except for Sole Proprietor; and (k) Any other document, other than the documents under **ITT 11.1 (a) – (j)** above,

			required to be completed and submitted by Tenderers, as specified in the **TDS.**
		11.2	Where a sample(s) is required by a PE, the sample shall be: (a) Physically submitted as part of the tender, in the quantities, sizes and other details requested in the **TDS**; (b) carriage paid; (c) received on, or before, the closing time and date for the submission of tenders and at the address shown in the **TDS**; and (d) Evaluated to determine compliance with all characteristics listed in the **TDS**.
		11.3	The PE shall reject the tender if the sample(s) does not conform to all characteristics prescribed in the tendering documents and are not submitted within specified time; and shall retain the sample(s) of the successful tenderer.
		11.4	Where it is not possible to avoid using a propriety article as a sample, a tenderer shall make it clear that the propriety article is displayed only as an example of the type or quality of the goods being tendered for and that competition shall not thereby be limited to that article only.
		11.5	Samples made up from materials supplied by a PE shall not be returned to a tenderer nor shall a PE be liable for the cost of making them.
		11.6	All samples produced from materials belonging to an unsuccessful tenderer which are not claimed by the tenderer within a period of thirty (30) days from the date of award of contract

			shall be the property of the PE and shall dispose them in such a manner as may be directed by the Accounting Officer.
12.	Documents Establishing Eligibility of Goods and Related Services and Conformity to Tendering Documents	12.1	Pursuant to **ITT 11 [Documents and Sample(s) Constituting the Tender]**, the Tenderer shall furnish, as part of its tender, documents establishing the eligibility of the Health Sector Goods and Related Services to be supplied under the Contract.
		12.2	The documentary evidence of the eligibility of the goods and related services shall consist of a statement in the Price Schedule of the country of origin of the goods and related services offered which shall be confirmed by a certificate of origin issued at the time of shipment.
		12.3	The documentary evidence of conformity of the Goods and Related Services to the Tendering Documents may be in the form of literature, drawings, and data and shall consist of: a) a detailed description of the essential technical and performance characteristics of the Goods; b) an item-by-item commentary on the PE's Technical Specifications demonstrating substantial responsiveness of the Goods and Related Services to those specifications, or a statement of deviations and exceptions to the provisions of the Technical Specifications; (c) any other procurement specific documentation requirement as stated in the **TDS.**
		12.4	(a) Unless the **TDS** stipulates otherwise, the Goods to be supplied under the contract shall be

			registered with the relevant authority in the United Republic of Tanzania. A Tenderer who has already registered its Goods by the time of Tendering should submit a copy of the Registration Certificates, with its Tender. Otherwise, the successful Tenderer, by the time of contract signing, shall submit to the PE either; (b) A copy of the Registration Certificate of the Goods for use in the United Republic of Tanzania, OR if such Registration Certificate has not yet been obtained; (c) Evidence establishing to the PE's satisfactions that the Tenderer has complied with all the documentary requirements for registration as specified above
		12.5	The PE shall at all times cooperate with the successful Tenderer to facilitate the registration process within the United Republic of Tanzania
		12.6	If the Goods of the successful Tender have not been registered in the United Republic of Tanzania at the time of contract signing, then the contract shall become effective upon such date as the certificate of Registration is obtained.
		12.7	For purposes of the commentary to be furnished pursuant to **ITT 12.3 (b)** above, the Tenderer shall note that standards as well as references to brand names designated by the PE in its Specifications are intended to be descriptive only and not restrictive. The Tenderer may substitute alternative standards, brand names, and/or catalogue numbers in its tender, provided that it demonstrates to the PE's satisfaction that the substitutions ensure substantial equivalence to those designated in the Specifications.

		12.8	The required documents and other accompanying documents must be in the language of the tender specified in **ITT 10 [Language of Tender]**. In case any other language than language of tender is used, the pertinent translation into that language of tender shall be attached to the original version.
13.	Documents Establishing Eligibility and Qualifications of the Tenderer	13.1	The Tenderer shall upload, as part of its Tender, documents establishing the Tenderer's eligibility to Tender and its qualifications to perform the contract if its Tender is accepted. In the event that pre-qualification of potential Tenderers has been undertaken, only tenders from pre-qualified Tenderers will be considered for award of contract. These qualified Tenderers should submit their Tenders with any information updating the original pre-qualification applications or, alternatively, confirm in their tenders that the originally submitted pre-qualification information remains essentially correct as of the date of tender submission. The update or confirmation should be provided in **Section V [Tendering Forms].**
		13.2	The documentary evidence of the Tenderer's eligibility to tender shall establish to the PE's satisfaction that the Tenderer, at the time of submission of its tender, is from an eligible country as defined under **ITT 4 [Eligible Goods and Related Services].**
		13.3	The documentary evidence of the Tenderer's qualifications to perform the contract if its Tender is accepted shall establish to the PE's satisfaction that: a) in the case of a Tenderer offering to supply

			goods under the contract which the Tenderer did not manufacture or otherwise produce, the Tenderer has been duly authorized by the goods' Manufacturer or producer to supply the goods in the United Republic of Tanzania; b) in the case of a Tenderer not doing business within the United Republic of Tanzania, the Tenderer is or will be (if awarded the contract) represented by an Agent in that country equipped, and able to carry out maintenance, repair, and spare parts stocking obligations prescribed in the General Conditions of Contract and/or Technical Specifications on behalf of the Supplier; and c) the Tenderer meets each of the qualification criterion specified in **Section IV [Qualification and Evaluation Criteria]**.
		13.4	Tenderers can get information on national standards and registration process from Regulators at the appropriate address specified in the **TDS**.
		13.5	Tenders submitted by a JVCA shall comply with the following requirements: a) the tender shall include all the information listed in pursuant to **ITT13.3** above for each JVCA partner; b) one of the partners of the JVCA will be nominated as being in charge, and this authorization shall be evidenced by submitting a Power of Attorney signed by legally authorized signatories of all the partners of the JVCA; c) a copy of the JVCA agreement entered into by all partners shall be submitted with the tender. Alternatively, a Letter of Intent to

			execute a JVCA agreement shall be signed by all partners and submitted with the tender, together with a copy of the proposed Agreement; d) the Tender Securing Declaration as stated in accordance with **ITT 20 [Tender Security or Tender Securing Declaration]**, and in case of a successful tender, the Agreement, shall be signed so as to be legally binding on all partners of the JVCA; and e) Any exception to the documents listed in (a) to (d) above will be indicated in **TDS**.
14.	Lot and Package-Multiple Contracts	14.1	Where a tender is divided into lots or packages, the number, nature, location and size of each lot or package will be as indicated in the **TDS**. The minimum and maximum number of lots or packages for which a tenderer may tender will be as indicated in the **TDS**.
		14.2	When tendering for more than one Contract under the lot and package arrangements, the Tenderer must provide evidence that it meets or exceeds the sum of all the individual requirements for the lots or package being applied for in regard to:- a) average annual turnover; b) particular experience; c) financial means; and d) personnel capabilities. as provided in **Section IV [Qualification and Evaluation Criteria]**.
		14.3	Incase the Tenderer fails to fully meet any of these criteria, it may be qualified only for those lots/multiple contracts for which the Tenderer

			meets the criteria as provided in **Section IV [Qualification and Evaluation Criteria]**.
		14.4	Each lot shall form a separate contract unless otherwise stated in the **TDS**.
		14.5	Where lots are to be apportioned to different tenderers, the PE may require the tenderer for a particular lot to ensure the coordination of the execution of the lots as specified in **TDS**.
15.	Form of Tender and Price Schedules	15.1	The Tenderer shall fill the Form of Tender furnished in the Tendering Documents through the system. The Form of Tender must be completed without any alterations to its format and no substitute shall be accepted.
16.	Alternative Tenders by Tenderers	16.1	Tenderers shall submit offers that comply with the requirements of the Tendering Documents, including the basic Tenderer's specification as indicated in the specifications and Schedule of Requirements. Alternatives will not be considered, unless specifically allowed for in the **TDS**. If so allowed, **ITT 16.2** shall prevail.
		16.2	When alternative schedule for delivery of goods is explicitly invited, a statement of that effect will be included in the **TDS** as will the method for evaluating different schedule for delivery of goods.
		16.3	If so allowed in the **TDS**, Tenderers wishing to offer technical alternatives to the requirements of the Tendering documents must also submit a Tender that complies with the requirements of the Tendering documents, including the basic technical design as indicated in the specifications. In addition to submitting the basic Tender, the

			Tenderer shall provide all information necessary for a complete evaluation of the alternative by the PE, including specifications, breakdown of prices, and other relevant details.
17.	Tender Prices and Discounts	17.1	The Tender prices and discounts quoted by the Tenderer in the Form of Tender and in the Price Schedules shall conform to the requirements specified in **ITT 17.1 to 17.11.**
		17.2	All items in the schedule of requirements must be listed and priced separately in the Price Schedules. If a Price Schedule shows item listed but not priced the Tender will be rejected as being substantially non-responsive. Items not listed in the Price Schedule shall be assumed to be not included in the Tender and theTender will be rejected as being substantially non-responsive.
		17.3	The tender price to be quoted in the Form of Tender in accordance with I**TT 15.1 [Form of Tender and Price Schedules]** shall be the total price of the tender, including any discounts offered.
		17.4	The Tenderer shall quote any unconditional discounts and the methodology for their application in the Form of Tender in accordance with **ITT 15.1 [Form of Tender and Price Schedule]**. Where a tender is divided in lots or packages,Tenderer may quote any discount and the methodology for their application in the Form of Tender in accordance with **ITT 15 [Form of Tender]** in the event of consolidation of some or all of the lots or packages for which he has submitted individual tenders.
		17.5	Prices quoted by the Tenderer shall be fixed

			during the Tenderer's performance of the Contract and not subject to variation on any account, unless otherwise specified in the **TDS**. A Tender submitted with an adjustable price quotation shall be treated as non responsive and shall be rejected, pursuant to ITT 28 [Preliminary Examination of Tenders]. However, if in accordance with the **TDS**, prices quoted by the Tenderer shall be subject to adjustment during the performance of the Contract, a Tender submitted with a fixed price quotation shall not be rejected, but the price adjustment shall be treated as zero.
		17.6	If so specified in ITT 1.1, Tenders are being invited for individual lots (contracts) or for any combination of lots (packages). Unless otherwise specified in the **TDS**, prices quoted shall correspond to 100 % of the items specified for each lot and to 100% of the quantities specified for each item of a lot. Tenderers wishing to offer discounts for the award of more than one Contract shall specify in their Tender the price reductions applicable to each package, or alternatively, to individual Contracts within the package. Discounts shall be submitted in accordance with ITT 17.4 provided the Tenders for all lots (contracts) are opened at the same time.
		17.7	The terms EXW, CIP, and other similar terms shall be governed by the rules prescribed in the current edition of Incoterms, published by The International Chamber of Commerce, as specified in the **TDS.**
		17.8	Prices shall be quoted as specified in each Price Schedule included in Section V, Tendering

			Forms. The dis-aggregation of price components is required solely for the purpose of facilitating the comparison of Tenders by the PE. This shall not in any way limit the PE's right to contract on any of the terms offered. In quoting prices, the Tenderer shall be free to use transportation through carriers registered in any eligible country, in accordance with Section V, Eligible Countries. Similarly, the Tenderer may obtain insurance services from any eligible country in accordance with Section VI, Eligible Countries. Prices shall be entered in the following manner: (a) for Goods manufactured in the United Republic of Tanzania: (i) the price of the Goods quoted EXW (ex-works, ex-factory, ex warehouse, ex showroom, or off-the-shelf, as applicable), including all customs duties and sales and other taxes already paid or payable on the components and raw material used in the manufacture or assembly of the Goods; (ii) any sales tax and other taxes which will be payable on the Goods if the Contract is awarded to the Tenderer; and (iii) the price for inland transportation, insurance, and other local services required to convey the Goods to their final destination (Project Site) specified in the **TDS**; (b) for Goods manufactured outside the United Republic of Tanzania, to be imported: (i) the price of the Goods, quoted CIP named place of destination, in the Purchaser's Country, as specified in the **TDS**; and (ii) the price for inland transportation, insurance, and other local services required to convey the

			Goods from the named place of destination to their final destination (Project Site) specified in the **TDS**; (c) for Goods manufactured outside the United Republic of Tanzania, already imported: (i) the price of the Goods, including the original import value of the Goods; plus any mark-up (or rebate); plus any other related local cost, and custom duties and other import taxes already paid or to be paid on the Goods already imported; (ii) the custom duties and other import taxes already paid (need to be supported with documentary evidence) or to be paid on the Goods already imported; (iii) the price of the Goods, obtained as the difference between (i) and (ii) above; (iv) any sales and other taxes which will be payable on the Goods if the Contract is awarded to the Tenderer; and (v) the price for inland transportation, insurance, and other local services required to convey the Goods from the named place of destination to their final destination (Project Site) specified in the **TDS**. (d) for Related Services, other than inland transportation and other services required to convey the Goods to their final destination, whenever such Related Services are specified in the Schedule of Requirements: the price of each item comprising the Related Services (inclusive of any applicable taxes).
		17.9	In the case of National, International and Restricted Competitive Tendering on Fixed

			Budget Method, the Tender Price quoted by the Tenderer shall not exceed the budget indicated in the **TDS**. Any Tenderer whose Tender Price exceeds the provided budget, shall be rejected.
18.	Tender Currencies	18.1	Prices shall be quoted in the following currencies: a) For goods and services that the Tenderer will supply from within the United Republic of Tanzania, the prices shall be quoted in Tanzania Shillings, unless otherwise specified in the **TDS**. b) For goods and related services that the Tender will supply from outside the United Republic of Tanzania, or for imported parts or components of goods and related services originating outside the United Republic of Tanzania, the Tender prices shall be quoted in any freely convertible currency of another country. If the Tenderer wishes to be paid in a combination of amounts in different currencies, it may quote its price accordingly but use no more than three foreign currencies
		18.2	Unless otherwise specified in the **TDS**, local transportation, insurance and other services incidental to delivery of the goods covered under **ITT 18.1** and installation services covered under **ITT 18.2** shall be quoted in either foreign and/or local currency, depending upon the currency in which the costs are to be incurred and in accordance with the provisions of **ITT 18.1 (a) and (b)** above.
		18.3	The rates of exchange to be used by the Tenderer in arriving at the local currency shall be the selling rates for similar transactions established by the authority specified in the **TDS** twenty-eight (28) days prior to the date specified

			for opening of tenders.
		18.4	Tenderers may be required by the PE to clarify their foreign currency requirements and to substantiate that the amounts included in Lump Sum and in the **SCC** are reasonable and responsive to **ITT 18.1**.
19.	Tender Validity Period	19.1	Tenders shall remain valid for the period stipulated in the **TDS** after the date of Tender submission specified in **ITT 23 [Deadline for Submission of Tenders]**. A Tender valid for a shorter period shall be rejected by the PE as non-responsive.
		19.2	In exceptional circumstances, prior to expiry of the original Tender validity period, the PE may request that the Tenderers consent to an extension of the period of validity of their Tenders. The request and the Tenderers responses shall be made through NeST.
		19.3	The Tender Security provided under **ITT 20 [Tender Security or Tender Securing Declaration]** shall also be suitably extended. A Tenderer may refuse the request without forfeiting its Tender Security or causing to be executed its Tender Securing Declaration.
		19.4	A Tenderer agreeing to the request will not be required or permitted to modify its Tender, but will be required to extend the validity of its Tender Security or Tender Securing Declaration for the period of the extension, and in compliance with **ITT 20 [Tender Security or Tender Securing Declaration]** in all respects.
		19.5	In the case of fixed price contracts, if the award is delayed by a period exceeding sixty (60) days

			beyond the expiry of the initial Tender validity period, the contract price may be adjusted by a factor specified in the request for extension.
20.	Tender Security or Tender Securing Declaration	20.1	Pursuant to **ITT 11 [Documents and Sample(s) Constituting the Tender]**, unless otherwise specified in the **TDS**, the Tenderer shall furnish as part of its tender, a Tender Security in original form and in the amount and currency specified in the TDS or Tender Securing Declaration as specified in the **TDS** in the format provided in **Section V [Tendering Forms].**
		20.2	The Tender security or Tender securing declaration is required to protect the PE against the risk of Tenderer's conduct which would warrant the security's forfeiture, pursuant to **ITT 20.9.**
		20.3	The Tender Security shall be denominated in local currency or in a freely convertible currency, and shall be, at the Tenderer's option, in one of the following forms: a) Unconditional bank guarantee, an irrevocable letter of credit issued by a reputable bank, or an insurance bond issued by a reputable insurance firm located in the United Republic of Tanzania or abroad valid for twenty-eight (28) days beyond the end of the validity of the Tender. This shall also apply if the period for Tender validity is extended. In either case, the form must include the complete name of the Tenderer; or (b) Another security if indicated in the **TDS**.
		20.4	The Tender Security shall be in accordance with the Form of the Tender Security or Tender

			Security Declaration included in Section V – Tendering Forms or another form approved by the PE prior to the Tender submission
		20.5	The Tender Security or Tender- Securing Declaration of a JVCA must be in the name of the JVCA that submits the tender. If the JVCA has not been legally constituted into a legally enforceable JVCA at the time of tendering, the tender security or Tender-Securing Declaration shall be in the names of all future members as named in the letter of intent referred to in ITT 13.5 and ITT 11.1 [Documents and Sample(s) Constituting the Tender]
		20.6	Any Tender not accompanied by a Tender security or Tender Securing Declaration in accordance with ITT 20.1 or 20.3 shall be rejected by the PE as non-responsive, pursuant to ITT 28 [Preliminary Examination of Tenders].
		20.7	Unsuccessful Tenderers' Tender Security will be discharged or returned as promptly as possible, but not later than thirty (30) days after the expiration of the period of Tender validity prescribed by the PE pursuant to **ITT 19 [Tender Validity Period]**. The PE shall make no claim to the amount of the tender security, and shall promptly return the tender security document, after whichever of the following that occurs earliest: (a) the expiry of the tender security; (b) the entry into force of a procurement contract and the provision of a security for the performance of the contract if such a security is required by the tender documents;

			(c) the rejection by the PE of all tenders; (d) the withdrawal of the tender prior to the deadline for the submission of tenders, unless the tender documents stipulate that no such withdrawal is permitted.
		20.8	The successful Tenderer's Tender Security will be discharged upon the Tenderer furnishing the performance security, pursuant to **ITT 43 [Performance Security]** and signing the contract pursuant to **ITT 44 [Signing of Contract]**.
		20.9	In case of Tender Security, it shall be forfeited if: a) a Tenderer withdraws its Tender during the period of Tender validity specified by the Tenderer on the Form of Tender except as provided for in **ITT 19.2 [Tender Validity Period]**; b) A successful Tenderer fails to furnish performance security in accordance with **ITT 43 [Performance Security]** or fails to sign the contract in accordance with **ITT 44 [Signing of Contract].**
		20.10	In the case of the Tender Securing Declaration, it shall be executed; a) if a Tenderer withdraws its tender during the period of Tender validity specified by the Tenderer on the Form of Tender except as provided for in **ITT 19.2 [Tender Validity Period]**; or b) in the case of a successful Tenderer, if the Tenderer fails to furnish performance security in

			accordance with **ITT 43 [Performance Security]** or fails to sign the contract in accordance with **ITT 44 [Signing of Contract]**.
		20.11	The failure of a Tenderer to abide with the terms of Tender Securing Declaration shall be reported to the Public Procurement Authority (PPRA) for debarment for a period which they shall determine. A Tenderer debarred by PPRA shall be ineligible to participate in during the period of debarment.
21.	Format and Signing of Tender	21.1	The Tenderer shall prepare documents comprising the tender as described in **ITT 11 [Documents and Sample(s) Constituting the Tender]**.
		21.2	TheTender shall be signed by a person or persons duly authorized to sign on behalf of the Tenderer except where the signatory of the tender is a Sole Proprietor. Incase of a person signing on behalf of the Tenderer, the authorization document(s) shall be a dully notarized Power of Attorney in the format provided in **Section V [Tendering Forms]** which must be submitted together with the tender indicating position of the signatory. A signatory who is a Sole Proprietor shall submit an Extract from Registrar issued by BRELA or any other document from competent Business line authority indicating name of business owner. The acceptable authorization document(s) is/are as listed in the **TDS**.
		21.3	The Tenderer shall furnish information as described in the Form of Tender on commissions or gratuities, if any, paid or to be

			paid to agents relating to this tender and to contract execution if the Tenderer is awarded the contract.

D. Submission of Tenders

22.	Submission of Tenders	22.1	Tenders submitted through NeST shall considered to be true and legal versions duly authorized and duly executed by the tenderer and intended to have binding legal effect.
		22.2	The tender shall bear e-signature or digital signatures, where applicable for identity and authentication purposes and the identity of the tenderer may be verified with a follow-up due diligence process.
		22.3	Tenders submitted through NeST shall be received in full prior to the closing time and the tenderers shall receive an acknowledgment of receipt of their tenders or amendment through the system.
		22.4	Tenderers must ensure the integrity, completeness, and authenticity of their submission; and in case of electronic records entered online and files containing the tender being unreadable for any reason, the tender submitted shall not be considered.
		22.5	In addition, if required in accordance with **ITT 11** [**Documents and Sample(s)** Constituting the Tender], the Tenderer shall deliver any samples at the address shown in the **TDS** not later than the deadline for submission of tenders.
23.	Deadline for Submission of Tenders	23.1	Tenders shall be received by the PE through NeST in a manner specified under **ITT 22.2 [Submission of Tenders]** not later than the date and time specified in the **TDS**.
		23.2	The PE may, in exceptional circumstances and at its discretion, extend the deadline for the submission of tenders by amending the

			Tendering documents in accordance with **ITT 9 [Amendment of Tender Documents**], in which case all rights and obligations of the PE and Tenderers previously subject to the deadline will thereafter be subject to the new deadline.
24.	Late Tenders	24.1	NeST does not allow a Tenderer to submit its tender after the deadline for submission of tenders in accordance with **ITT 23** [**Deadline for Submission of Tenders**].
25.	Modification, Substitution, and Withdrawal of Tenders	25.1	A Tenderer may modify or substitute or withdraw its Tender after it has been submitted to the PE through NeST. Such modification or substitution or withdrawal should be made prior to the deadline for submission of Tenders. The tenderer shall receive an acknowledgment of receipt of any amendment of its submitted tender through the system.
		25.2	No Tender may be withdrawn or replaced or modified in the interval between the deadline for submission of Tenders and the expiration of the period of Tender validity specified by the Tenderer on the Form of Tender. Withdrawal of a Tender during this interval shall result in execution of Tender Securing Declaration, pursuant to the **ITT 20.10** [**Tender Security or Tender Securing Declaration**].

E. Opening and Evaluation of Tenders

26.	Opening of Tenders	26.1	The opening shall be done through NeST as per time specified in the tender documents. The Tender opening records shall be made available through appropriate sections of NeST.
		26.2	A Tenderer or any other person with interest in the tender process can access tender opening records on the appropriate section of NeST.
27.	Confidentiality	27.1	Information relating to the examination, clarification, evaluation, and comparison of tenders and recommendation of contract award shall not be disclosed to Tenderers or any other persons not officially concerned with such process until the notice of intention to award the contract to the successful Tenderer has been issued.
		27.2	Any effort by a Tenderer to influence the PE's processing of Tenders or award decisions may result in the rejection of his Tender.
28.	Clarification of Tenders	28.1	In order to assist in the examination, evaluation, and comparison of Tenders and post-qualification of the Tenderers, the PE may, at its discretion, ask any Tenderer for a clarification of its Tender including a breakdown of prices. Any clarification submitted by a Tenderer that is not in response to a request by the PE shall not be considered.
		28.2	The request for clarification and the response shall be communicated through NeST and The Tenderer shall; respond through NeST.

			No change in the prices or substance of the Tender shall be sought, offered, or permitted except provided otherwise.
29 .	Preliminary Examination of Tenders	29.1	Prior to the detailed evaluation of Tenders, the PE will determine whether each Tender: a) meets the eligibility criteria defined in **ITT 3 [Eligible Tenderers]** and **ITT4 [Eligible Goods and Related Services]**; b) has been properly signed; c) is accompanied by the required securities; and d) is substantially responsive to the requirements of the Tendering Documents. The PE's determination of a Tender's responsiveness will be based on the contents of the Tender itself.
		29.2	A substantially responsive Tender is one which conforms to all the terms, conditions, and specifications of the Tendering Documents, without material deviation, omission or reservation. A material deviation or reservation is one that: - a) if accepted, would affect in any substantial way the scope, quality, or performance of the Goods and Related Services in the Contract; or limit in any substantial way, inconsistent with the Tendering Documents, the PE's rights or the Tenderers obligations under the Contract; or b) if rectified, would unfairly affect the competitive position of other Tenderers presenting substantially responsive Tenders. For the purpose of this section, the following

			definitions apply; "**Deviation**" is a departure from the requirements specified in the Tendering Document; "**Omission**" is the failure to submit part or all of the information or documentation required in the Tendering Document; and "**Reservation**" is the setting of limiting conditions or withholding from complete acceptance of the requirements specified in the Tendering Document.
		29.3	The PE will confirm that the documents and information specified under **ITT 11 [Documents and Samples Constituting the Tender], ITT 12** [**Documents Establishing Eligibility of Goods and Related Services and Conformity to Tendering Documents]** and **ITT 13 [Documents Establishing Eligibility and Qualification of a Tenderer]** have been provided in the Tender. If any of these documents or information is missing or is not provided in accordance with the Instructions to Tenderers, the Tender shall be rejected.
		29.4	The PE may waive any minor informality, nonconformity, or irregularity in a Tender which does not constitute a material deviation, provided such waiver does not prejudice or affect the relative ranking of any Tenderer.
		29.5	Provided that a tender is substantially responsive, the PE may request that the Tenderer submit the necessary information or documentation, within a reasonable period of

			time, to rectify non-material non-conformities or omissions in the tender related to documentation requirements. Such omission shall not be related to any aspect of the price of the tender. Failure of the Tenderer to comply with the request may result in the rejection of its tender.
		29.6	Provided that a tender is substantially responsive, the PE shall rectify quantifiable non-material and non- conformities related to the Tender Price. To this effect, the Tender Price shall be adjusted, for comparison purposes only, to reflect the price of a missing or non- conforming item or component.
		29.7	If a Tender is not substantially responsive, to the requirements of the Tendering Documents, it shall be rejected by the PE and may not subsequently be made responsive by the Tenderer by correction of the material deviation, reservation or omission.
		29.8	Material deviations to commercial terms and conditions, which justify rejection of a tender shall include the following: a) failure to sign the Form of Tender and price schedules by the authorized person or persons; b) failure to satisfy eligibility requirements; c) failure to submit a tender security as specified in the tendering documents; d) failure to satisfy the tender validity period; e) inability to meet the critical delivery schedule specified in the tendering

			documents, where such schedule is a crucial condition with which tenderers must comply; f) failure to comply with minimum experience criteria as specified in the tendering documents; g) conditional tenders such as conditions in a tender which limit the tenderer's responsibility to accept an award; h) inability to accept the price adjustment formulae of the tendering documents; i) stipulating price adjustment when fixed price tenders were invited; j) subcontracting in a substantially different amount or manner than that permitted; and k) failure to submit major supporting documents required by the tendering documents to determine substantial responsiveness of a tender.
		29.9	All tenders shall be checked for substantial responsiveness to the technical requirements of the tendering documents and non-conformity to technical requirements, which are justifiable grounds for rejection of a tender includes the following: a) failure to tender for the required scope of tender as instructed in the tendering documents and where failure to do so has been indicated as unacceptable; b) failure to quote for a major item in the package; c) failure to meet major technical requirements, such as offering completely different types of equipment or materials from the types specified, plant capacity well

			below the minimum specified, equipment not able to perform the basic functions for which it is intended; and d) presentation of absolutely unrealistic and inadequate implementation plans and schedules regarding performance, technical or service factors.
30.	Conversion to Single Currency	30.1	To facilitate evaluation and comparison of tenders, the PE will convert all Tender prices expressed in the amounts in various currencies in which the Tender prices are payable in Tanzania Shillings at the selling exchange rate established for similar transactions by the BOT at the date of tender opening.
		30.2	The currency selected for converting Tender prices to a common base for the purpose of evaluation, along with the source and date of the exchange rate, are specified in the **TDS**.
31.	Detailed Evaluation and Comparison of Tenders	31.1	The PE shall evaluate and compare only the Tenders determined to be substantially responsive, pursuant to **ITT 29 [Preliminary Examination of Tenders]**.
		31.2	To evaluate a Tender, the PE shall consider the following: a) evaluation will be done for Items or Lots (contracts), as specified in the **TDS**; and the Tender Price as quoted in accordance with **Clause 14** [Form of Tender and Price Schedules]*;* b) price adjustment due to discounts offered in accordance with **ITT 17.4 [Tender Prices and Discounts]**;

			c) converting the amount resulting from applying (a) if relevant, to a single currency in accordance with **ITT 30 [Conversion to Single Currency]**; d) price adjustment due to quantifiable non-material non-conformities in accordance with **ITT 29.5 and 29.6 [Preliminary Examination of Tenders]**; and e) the additional evaluation factors are specified in **Section IV [Qualification and Evaluation Criteria]**.
		31.3	The PE reserves the right to accept or reject any variation, deviation, or alternative offer. Variations, deviations, and alternative offers and other factors which are in excess of the requirements of the Tendering Documents or otherwise result in unsolicited benefits for the PE will not be taken into account in Tender evaluation.
		31.4	The estimated effect of the price adjustment provisions of the Conditions of Contract, applied over the period of execution of the Contract, shall not be taken into account in tender evaluation
		31.5	If these Tendering Documents allows Tenderers to quote separate prices for different lots (contracts), the methodology to determine the lowest evaluated price of the lot (contract) combinations, including any discounts offered in the Form of Tender, is specified in **Section IV [Qualification and Evaluation Criteria]**.
		31.6	The PE's evaluation of a Tender will exclude and not take into account:

			a) in the case of goods manufactured in the United Republic of Tanzania or goods of foreign origin already located in the United Republic of Tanzania, sales and other similar taxes, which will be payable on the goods if a contract is awarded to the Tenderer; b) in the case of goods of foreign origin offered from abroad, customs duties and other similar import taxes which will be payable on the goods if the contract is awarded to the Tenderer; and c) any allowance for price adjustment during the period of execution of the contract, if provided in the Tender.
		31.7	The PE's evaluation of a tender may require the consideration of other factors, in addition to the Tender Price quoted in accordance with **ITT 17 [Tender Prices and Discounts]**. These factors may be related to the characteristics, performance, and terms and conditions of purchase of the Goods and Related Services. The effect of the factors selected, if any, shall be expressed in monetary terms to facilitate comparison of tenders, unless otherwise specified in the **TDS** from amongst those set out in **Section IV [Qualification and Evaluation Criteria]**. The criteria and methodologies to be used shall be as specified in ITT 31.2 (e).
		31.8	The comparison shall be between the EXW price of the goods offered from within the United Republic of Tanzania, such price to include all costs, as well as duties and taxes paid or payable on components and raw material incorporated or to be incorporated in

			the goods, and the CIF named port of destination (or CIP border point, or CIP named place of destination) price of the goods offered from outside the United Republic of Tanzania.
32.	National Preferences	32.1	Any procurement of goods utilizing this Standard Tendering Document shall provide local tenderers with a margin of preference as indicated in **Section IV [Qualification and Evaluation Criteria]** during tender evaluation.
		32.2	To benefit with national preference schemes, a local tenderer shall meet the following criteria: - a) for individual company:- (i) Should be incorporated or registered inthe United Republic. (ii) The majority of the paid-up share capital of the company is owned either by the Government or by citizens of Tanzania. The tenderer shall submit the Shareholding Structure of the company . (iii) That no arrangement of any major part of the net profits or other tangible benefits of the domestic company will accrue or be paid to persons who are not citizens of Tanzania or to companies which would not be eligible for preference. b) for Joint Ventures of local company- (i) individual member companies are incorporated or registered in the United Republic. (ii) majority of the paid-up share capital of the individual companies are held by citizens

			of Tanzania; (iii) the Joint Venture itself is registered in the United Republic; or (iv) do not have arrangement whereby any major part of the net profits will accrue or be paid to persons who are not citizens of Tanzania or to companies which would not be eligible for preference. c) for partners or individual persons trading as supplier, the majority of shares shall be held by citizens of Tanzania. For the purpose of **ITT 34.2** Individual company, Individual company forming the JV, JVs of local company, partners and individual should satisfy the requirements specified under **Section IV [Qualification and Evaluation Criteria]**.
		32.3	Where a JV is formed between a local firm and a foreign firm for purposes of qualifying for national preference, at least 51% of the paid-up share capital of the local firm shall be owned by the Government or citizens of the Tanzania.
		32.4	Where a JV is formed between a local firm and a foreign firm for purposes of qualifying for national preference, the local firm to the JV shall allocate not less than 50% of key personnel required for the implementation of the project and the JV Agreement shall indicate the titles of key personnel that will be contributed by each party to the JV.
		32.5	The contribution of the financial resources of the local firm to the JV shall not be less than 50% of the financial resources required to

			implement the contract.
		32.6	For a tenderer to benefit from the preference for local goods, shall indicate in its tender the intent to use goods domestically produced, mined or manufactured in the United Republic during implementation of the contract.
		32.7	Subject to **ITT 32.6,** a PE shall grant a preference of 15%, its application and detail of evaluation shall be specified in **Section IV [Qualification and Evaluation Criteria]**.
33.	Exclusive Preference	33.1	In the event a foreign tenderer is invited to participate in tenders under exclusive preference as indicated in Section IV [Qualification and Evaluation Criteria] TDS, it shall enter into a JV or sub-contract with a local firm and the local firm shall meet the eligibility criteria specified under ITT 32.2 (a).
		33.2	In case of JV for tenders set aside for exclusive preference, local firm contribution to the JV shall not be less than 75%. The JV shall submit a Letter of Intent to form Joint Venture (JV) accompanied with a draft JV Agreement or a Joint Venture Agreement showing the percentage of the contribution.
		33.3	In case of sub-contracting for tenders set aside for exclusive preference, the foreign firm shall subcontract between 5% and 30% of the total contract value to a local firm. The foreign tenderer shall submit Letter of Intent to Sub-contract indicating percentage, type of works to be sub-contracted and the applicable method of sub-contracting

			between domestic or nominated. The works to be sub-contracted shall exclude Provisional Sums and Prime Cost.
		33.4	Details of evaluation for JV and Sub-contracting shall be specified in Section IV [Qualification and Evaluation Criteria].
34.	Preference to Partnerships	34.1	Tenderers may, for purposes of enjoying a margin of preference as indicated in Section IV [Qualification and Evaluation Criteria], during supply of goods, form a partnership between a local firm and local firm or a local firm and a foreign firm.
		34.2	A foreign firm shall receive preference if it forms a partnership with a local firm in which the local firm owns at least 51% of paid-up share capital (contribution) in the partnership and contributes at least 50% of key personnel in the implementation of the contract.
		34.3	The financial resources contribution of the local firm to the Partnership shall not be less than 50% of the financial resources required to implement the contract.
		34.4	PE shall grant a preference of 10% to a foreign firm that forms a partnership with local firm.
		34.5	The application of preference and details of evaluation is as specified in Section IV [Qualification and Evaluation Criteria].
35.	Preference to Local Suppliers Sub-	35.1	Local Supplier whose investment capital falls under the medium or large enterprises category, as classified in the Small and

	contracting to other Suppliers		Medium Enterprises National Development Policy of 2003 (as amended), may enjoy preference in supply tenders set aside for exclusive preference, provided they subcontract to a local Supplier whose investment capital falls under either micro or small enterprises category as classified in the Policy.
		35.2	Subject to ITT 35.1, a PE shall grant a margin of preference of up to 10%. Its application and detail of evaluation shall be specified in Section IV [Qualification and Evaluation Criteria].
		35.3	Local Supplier whose investment capital falls under the medium or large enterprises category shall submit a Letter of Intent to Sub-contract indicating percentage, type of goods for supply to be sub-contracted and method of sub-contracting.
36.	Determination of the Lowest Evaluated Tender	36.1	The Tender with the lowest evaluated price, from among those which are eligible, compliant and substantially responsive shall be the lowest evaluated Tender.
		36.2	In determining the lowered evaluated tenderer, the PE shall determine to its satisfaction whether the Tenderer that is selected as having submitted the lowest evaluated cost and substantially responsive Tender is eligible and meets the qualifying criteria specified in **ITT 11.1 [Documents and Samples Constituting the Tender]** as applicable, and **Section IV [Qualification and Evaluation Criteria]**.
		36.3	The determination shall be based upon an

			examination of the documentary evidence of the Tenderer's qualifications submitted by the Tenderer, pursuant to **ITT 13 [Documents Establishing Eligibility and Qualifications of the Tenderer]**. The determination shall not take into consideration the qualifications of other firms such as the Tenderer's subsidiaries, parent entities, affiliates, subcontractors or any other firm(s) different from the Tenderer.
		36.4	An affirmative determination shall be a prerequisite for award of the Contract to the Tenderer. A negative determination shall result in disqualification of the Tender, in which event the PE shall proceed to the Tenderer who offers a substantially responsive Tender with the next lowest evaluated cost to make a similar determination of that Tenderer's qualifications to perform satisfactorily.
		36.5	In the case of National, International and Restricted Competitive Tendering on Fixed Budget Method, the lowest evaluated price shall not exceed the provided budget. If the price of the Lowest Evaluated Tender exceeds the provided budget, it shall be rejected.
37.	Post-qualification of Tenderer	37.1	If pre- qualification was not undertaken, the Purchaser shall carry out the post-qualification of the Tenderer using only the requirements specified in **Section IV [Qualification and Evaluation Criteria]**.
		37.2	In the case of National, International and Restricted Competitive Tendering onFixed

			Budget Method, the lowest evaluated price shall not exceed the provided budget. If the price of the Lowest Evaluated Tender exceeds the provided budget, it shall be rejected.
		37.3	Where the tender price of the lowest evaluated Tenderer is considered to be abnormally low, the PE shall perform price analysis. Abnormally low tender means, in the light of the PE's estimate and of all the tenders submitted, the tender appears to be abnormally low by not providing a margin for normal levels of profit.
		37.4	The Purchaser may determine that a tender price is abnormally low by using several approaches, if possible: (i) Comparing the tender price with the pre-tender estimates. (ii) Comparing the tender price with the average tenders price offered by other tenderers submitting substantially responsive tenders. (iii) Comparing the tender price with prices paid in similar contracts in the recent past, either government or development partner-funded projects; (iv) Comparing the tender price with the available price cap set by Authority for specific assignment. The rate of determination for abnormally low tender will be as specified in the **TDS.**
		37.5	The following process shall apply: a) The Purchaser may reject a tender if the

			PE has determined that the price in combination with other constituent elements of the tender is abnormally low in relation to the subject matter of the procurement (scope of works or services) and raises concerns as to the ability of the Tenderer that presented that tender to perform the contract. b) Before rejecting an abnormally low tender, the PE shall request the Tenderer for an explanation of the tender or of those parts which it considers contribute to the tender being abnormally low; take account of the evidence provided in response to a request in writing; and subsequently verify the tender or parts of the tender being abnormally low. c) The decision of the PE to reject a tender and reasons for the decision shall be recorded in the procurement proceedings and promptly communicated to the Tenderer concerned. d) The PE shall not incur liability solely by rejecting abnormally low tender.
		37.6	The PE will determine to its satisfaction whether the Tenderer that is selected as having submitted the lowest evaluated responsive Tender is qualified to perform the contract satisfactorily, in accordance with the criteria listed in **Section IV [Qualification and Evaluation Criteria]**.
		37.7	The determination will take into account the Tenderer's financial, technical, and production capabilities as specified in **Section IV [Qualification and Evaluation**

			Criteria]. It will be based upon an examination of the documentary evidence of the Tenderer's qualifications submitted by the Tenderer, pursuant to **Section IV [Qualification and Evaluation Criteria]**, as well as such other information as the PE deems necessary and appropriate. Factors not included in these Tendering Documents shall not be used in the evaluation of the Tenderers' qualifications.
		37.8	A PE may seek independent references of a tenderer and the results of reference checks may be used in determining award of contract.
		37.9	In case of a foreign company, a PE shall seek independent reference of legal existence of a tenderer from Tanzania diplomatic missions abroad or from any other reliable source.
		37.10	An affirmative determination will be a prerequisite for award of the contract to the Tenderer. A negative determination will result in rejection of theTenderer's Tender, in which event the PE will proceed to the next lowest evaluated Tender to make a similar determination of that Tenderer's capabilities to perform satisfactorily.

F. Award of Contract

38.	**Criteria for Award**	38.1	Subject to ITT 36 [Determination of Lowest Evaluated Tender] and ITT 39 [Negotiations] the PE will award the Contract to the Tenderer whose Tender has been determined to be substantially responsive to the Tendering Documents and who has offered the lowest Evaluated Tender Price, provided that such Tenderer has been determined to be: a) eligible in accordance with the provisions of ITT 3 [Eligible Tenderers]; b) qualified to perform the Contract satisfactorily; and c) successful negotiations have been concluded, if any.
		38.2	If this Contract is being let on lots basis, the lowest evaluated Tender Price will be determined when evaluating this Contract in conjunction with other Contracts to be awarded concurrently, taking into account any discounts offered by the Tenderers for award of more than one Contract.
39.	Negotiations	39.1	Negotiations may be undertaken with the lowest evaluated Tender relating to the following areas: a) a minor alteration to the technical details of the statement of requirements or specifications; b) reduction of quantities for budgetary reasons, where the reduction is in excess of any provided for in the tendering documents; c) a minor amendment to the SCC; d) finalizing payment arrangements; e) delivery arrangements; f) the methodology or staffing; g) clarifying details that were not apparent or could not be finalized at the time of tendering; or h) reduction of Tender Price to match the available PEs Estimate and commensurate with the market prices, provided such reduction shall not make the tender abnormally low in

			accordance with ITT37.2 [Post-qualification of Tenderers]. Negotiation of price shall not be applicable for tenders invited under the National, International and Restricted Competitive Tendering on Fixed Budget method.
		39.2	Where negotiation fails to result into an agreement, the PE may invite the next ranked Tenderer for negotiations. Where negotiations are commenced with the next ranked Tenderer, the PE shall not reopen earlier negotiations.
		39.3	The results of any approved negotiations shall be specified in a letter of acceptance, incorporated into the appropriate contract document and the Minutes of Negotiations attached to the contract.
40.	PE's Right to Accept any Tender and to Reject any or All Tenders	40.1	Notwithstanding ITT 38 [Criteria for Award], the PE reserves the right to accept or reject any tender, and to annul the tendering process and reject all tenders at any time prior to the signing of the contract, without thereby incurring any liability to the affected Tenderer(s).
		40.2	Within seven (7) days from the date of rejection of Tenders in accordance with ITT 40.1 [PE's Right to Accept any Tender and to Reject any or All Tenders], Notice of the rejection of all tenders shall be given promptly through NeST to all Tenderers who have submitted Tenders.
		40.3	The PE shall upon request from any Tenderer communicate the ground(s) for rejection of the tender(s) but shall not be obliged to justify the given ground(s).
41.	PE's Right to Vary Quantities at the Time	41.1	The PE reserves the right at the time of contract award to increase or decrease the quantity of goods or related services originally specified in these Tendering documents (schedule of requirements) provided this does not exceed by the percentage indicated in the TDS, without any change in unit price or other terms and conditions of the Tender and Tendering Documents.

	of Award		
42.	Notification of Intention and Notification for Award	42.1	Prior to awarding of the contract, the PE shall issue a notice of intention to award the contract, in the format provided in Section IX [Contract Forms - Letter of Intention to Award the Contract],. the Notice shall be issued through NeST to all tenderers who participated in the tender in question giving them five (5) working days within which to submit complaint (s) to the PE thereof, if any.
		42.2	The condition stated above shall not apply where only one Tender has been received by PE in competitive method or where single source, national shopping, Minor value procurement method has been used.
		42.3	Where no complaints have been lodged, the Tenderer whose tender has been accepted will be notified by Letter of Acceptance in the format provided in Section IX [Contract Forms- Letter of Acceptance] through NeST, of the award by the PE prior to expiration of the Tender validity period.
		42.4	The notification of award will be part of the document forming the Contract, subject to the Tenderer furnishing the Performance Securing Declaration or Performance Security in accordance with ITT 43 [Performance Security/Performance Securing Declaration] and signing of the contract in accordance with ITT 44 [Signing of Contract].
		42.5	Upon the successful Tenderer's furnishing of the performance security or Performance Securing Declaration pursuant to ITT 43 [Performance Security or Performance Securing Declaration], the PE will promptly notify unsuccessful Tenderers, the name of the winning Tenderer and the Contract amount and will discharge the Tender Securing Declaration of the unsuccessful Tenderers pursuant to ITT 20.7 [Tender Security or Tender Securing Declaration].
43.	Perfor	43.1	Within fourteen (14) days after receipt of the Letter of

	mance Security/Performance Securing Declaration		Acceptance from the PE, and before signing of the contract, the successful Tenderer shall deliver to the PE a Performance Security or Performance Securing Declaration in the amount and in the form stipulated in the TDS and SCC, denominated in the type and proportions of currencies in the Letter of Acceptance and in accordance with the Conditions of Contract.
		43.2	Where applicable as indicated in the TDS, Within fourteen (14) working days after receipt of the Letter of Acceptance from the PE and before signing of the contract, the successful Tenderer shall deliver to the PE the Environmental and Social (ES) Performance Securing Declaration or Environmental and Social (ES) Performance Security in the amount and in the form stipulated in the TDS and SCC, denominated in the type and proportions of currencies in the Letter of Acceptance and in accordance with the Conditions of Contract.
		43.3	In the case of Performance Security, it shall be in the form specified in the TDS and SCC, and shall be in any of the following: (a) unconditional bank guarantee confirmed by a reputable local bank or, in the case of a successful foreign tenderer, bonded by a foreign bank; or (b) surety bond issued by any reputable surety or insurance company registered in the United Republic. Any Performance Security submitted shall be enforceable in the United Republic of Tanzania.
		43.4	In the case of Performance Securing Declaration, the successful Tenderer shall complete and submit a duly signed Declaration in the format provided in Section X [Contract Forms: Performance Securing Declaration].
		43.5	Failure of the successful Tenderer to comply with the requirement of ITT 39.1 shall constitute sufficient grounds for cancellation of the award and forfeiture of the Tender security or execution of the Tender Securing Declaration and any other

			remedies the PE may resort to awarding the Contract to the next ranked Tenderer or call for new tenders.
44.	Signing of Contract	44.1	Promptly after notification of award, PE shall send the successful Tenderer a draft Contract, incorporating all terms and conditions as agreed by the parties to the contract.
		44.2	Within fourteen (14) working days after furnishing the Performance Securing Declaration or Performance Security, the successful Tenderer and the PE shall sign the contract.
		44.3	Upon parties signing the Contract, the PE will promptly notify each unsuccessful Tenderer, the name of the successful Tenderer and the Contract amount and will discharge the Tender security of the Tenderers pursuant to ITT 20.7 [Tender Security or Tender Securing Declaration].
45.	Advance Payment	45.1	The PE will provide an Advance Payment on the Contract Price as stipulated in Conditions of Contract, subject to a maximum amount as stated in the TDS.
		45.2	The Advance Payment request shall be accompanied by an Advance Payment Security (Unconditional Bank Guarantee) and application form in the format provided in Section X [Contract Forms].
		45.3	For the purpose of receiving the Advance Payment, the Tenderer shall make and estimate of, and include in its tender, the expenses that will be incurred in order to commence Delivery of Goods.
46.	Fraudulent, Corrupt, Coercive, Collusive or	46.1	The PEs and Tenderers are required to observe the highest standard of ethics during the procurement and execution of such contracts. For the purpose of this provision, the following defined terms shall apply: - a) “coercive practice” means impairing or harming, or threatening to impair or harm directly or indirectly, any party or the property of the party for the purpose of influencing improperly the action or that party in connection with public

	Obstructive Practices		procurement or in furtherance of corrupt practice or fraudulent practice; b) "collusive practices" means impairing or harming, or threatening to impair or harm directly or indirectly, any part or the property of the Party for the purpose of influencing improperly the action or a part or in connection with public procurement or government contracting or in furtherance of a corrupt practice or a fraudulent practice; c) "**corrupt practice**" means the offering, giving receiving or soliciting of anything of value to influence the action of a public officer in the procurement process or contract execution; d) "fraudulent practice" means a misrepresentation of facts in order to influence a procurement process or the execution of a contract to the detriment of the Government or a public body and includes collusive practices among tenderers, prior to or after submission designed to establish tender prices at artificial non-competitive levels and to deprive the Government of the benefits of free and open competition; and e) "obstructive practice" means acts intended to materially impede access to required information in exercising a duty under the Act.
		46.2	The PE will reject a proposal for award of contract if it determines that the Tenderer recommended for award has engaged in corrupt, fraudulent, coercive, collusive and obstructive practices in competing for the contract.
		46.3	In pursuit of this policy the Government will cancel the portion of the funds allocated to a contract for goods, works, or services if it at any time determines that corrupt, coercive, fraudulent obstructive or fraudulent practices were engaged in by representatives of the PE or approving authority or of a beneficiary of the funds during the procurement or the execution of that contract, without the PE or approving authority having taken timely and appropriate action satisfactory to the Government of Tanzania to remedy the

			situation.
		46.4	A Tenderer will be debarred by the Public Procurement Regulatory Authority for a period of ten (10) years from participating in public procurement if at any time it determines that the firm has engaged in corrupt or fraudulent practices in competing for, or in executing, a public – financed contract.
		46.5	The Authority reserves the right, where a firm has been found by a foreign country, international organization or other foreign organization to have engaged incorrupt, coercive, collusive, fraudulent or obstructive practices, to declare that such a firm is ineligible, for a period as such firm(s) is debarred by the foreign country, international organization or other foreign organization to be awarded a public financed Contract in the United Republic of Tanzania.
		46.6	The Government of Tanzania will have the right to require that, in contract financed by the Government of Tanzania a provision be included requiring Tenderers to permit the Government of Tanzania to inspect their accounts and records relating to the performance of the contract and to have them audited by auditors appointed by the Government of Tanzania.
		46.7	Any communications between the Tenderer and the PE related to matters of alleged corrupt, coercive, collusive, fraudulent or obstructive practices must be made in writing or in electronic forms that provide record of the content of communication.

G. Review of Procurement Decisions

47.	Right to Review	47.1	A Tenderer who claims to have suffered or that may suffer any loss or injury as a result of a breach of a duty imposed on a PE or an approving authority in the course of these procurement proceedings may seek a review in accordance with the procedures set out hereunder.
48.	Time Limit on Review	48.1	The Tenderer shall submit an application for review within seven (7) working days of him becoming or should have become aware of the circumstances giving rise to the complaint or dispute.
49.	Submission of Applications for Review	49.1	Any application for administrative review shall be submitted through NeST to the Accounting Officer and a copy shall be electronically served to the PPRA.
		49.2	For PEs with delegated Procurement functions, applications for administrative review for tenders floated by the delegated Accounting Officer shall be submitted through NeST to the Accounting Officer with a copy served to delegated Accounting Officer and the Authority.
		49.3	The application for administrative review shall include: a) details of the procurement requirements to which the complaint relates; b) details of the provisions of the Act, Regulation or provision that has been breached or omitted; c) an explanation of how the provisions of the Act, Regulation, or provision has been breached or omitted, including the dates and name of the responsible public officer, where known; d) documentary or other evidence supporting the complaint where available; e) remedies sought; and f) any other information relevant to the complaint.

		49.4	Upon receipt of a complaint, the Accounting Officer shall suspend the procurement proceedings and notify all tenderers who participated in the tender within one(1) day after receipt of a complaint or dispute.
		49.5	The Accounting Officer (AO) shall not entertain a complaint or dispute or continue to do so after the procurement contract has entered into force.
		49.6	On receiving a complaint, the Accounting Officer may, depending on the nature of the complaint, constitute an independent review panel from within or outside his entity which shall review the complaint and advise him on the appropriate actions to be taken.
		49.7	Where the accounting officer constitutes an independent review panel, he shall notify all tenderers within one (1) day after receipt of a complaint or dispute.
50.	Decision by the Accounting Officer (AO)	50.1	The Accounting Officer shall, within five (5) working days after the receipt of the complaint/dispute and where an independent review panel is not constituted, or within seven (7) working days after the receipt of the complaint/dispute if an independent review panel is constituted, deliver a written decision which shall indicate: a) whether the application is upheld in whole, in part or rejected; b) the reasons for the decision; and c) any corrective measures to be taken.
		50.2	Where the Accounting officer of a PE does not issue a decision within the time specified in **ITT 50.1,** the Tenderer submitting the complaint or dispute or the PE shall be entitled immediately thereafter to institute proceedings under **ITT 47.1** [Review by the Public Procurement Appeals Authority] within five (5) working days after such specified time and upon instituting such

			proceedings, the competence of the Accounting Officer to entertain the complaint or dispute shall cease.
51.	Review by the Public Procurement Appeals Authority	51.1	Complaints or disputes which: (a) are not settled within the specified period under **ITT 50 [Decision by the Accounting Officer**]; (b) the tenderer is not satisfied with the decision of the accounting officer; or (c) arise after the procurement contract has entered into force pursuant to **ITT 44 [Signing of Contract]**, shall be referred to the Appeals Authority within five (5) working days from the date when the Tenderer received the decision of the accounting officer or, in case no decision is issued after the expiry of the time stipulated under **ITT 50.1** or when the Tenderer become aware or ought to have become aware of the circumstances giving rise to the complaint or dispute pursuant to **ITT 48.1 [Time Limit on Review]**.
		51.2	The Appeals Authority shall, within forty (40) days issue a written decision concerning the complaint or dispute stating the reasons for the decisions and the remedies granted if any.
		51.3	The decision of the Appeals Authority shall be binding to the parties on complaint, dispute or appeal and such decision shall be enforceable in the same manner as a decree or order of the court.
		51.4	Complaints/Appeals to PPAA shall be submitted through NeST at:https://nest.go.tz/module/complaint management/dashboard. The Physical Address for PPAA is as provided under the **TDS** .

SECTION III: TENDER DATA SHEET (TDS)

A. Introduction

S/N	Required Information/ Data	ITT Clause	ITT Clause Description
		A. Introduction	
1.	Name of Procuring Entity	1.1	MBEYA REGIONAL SECRETARIAT
2.	Subject of Procurement	1.1	**Procurement of medicines from prime vendor for Councils health facilities**
3.	Period of Supply of Goods	1.1	14 Days
4.	Duration of Contract	1.1	1095 Days
5.	Method of Procurement	1.2	National Competitive Tendering
6.	Financial Year	2.1	2025/2026
7.	Nature of the Supplies	2.1	Procurement of medicines from prime vendor for Councils health facilities
8.	Financing Institution	2.1 & 2.2	Not Applicable.
9	Name	2.1	Tender Number: 78/2025/2026/G/72

.	and Identification Number of the Contract		Tender Description: Procurement of medicines from prime vendor for Councils health facilities
10.	Eligible Tenderers	3.1	GOVERNMENT_ENTERPRISE, Company Local, Company Foreign, Special Group, Manufacturer Local, Sole Proprietor Local, Partnership Local, Manufacturer Foreign, Partnership Foreign and Sole Proprietor Foreign
11.	Number of JVCA	3.3	Not Applicable.
12.	Manufacturer's Authorization	4.6	**NOT_APPLICABLE**

B. Preparation of Tenderers

13	Language of the Tender	10.1	English
14	Samples requirements	11. 1(b)	The sample is not required
15	Additional Documents to be Submitted with the tender	11.1(i)	Not Applicable.
16	Address to submit sample	11.2(c)	Not Applicable.
17	Sample Characteristic	11.2(d)	Not Applicable
18	Other Procurement Specific Documentation evidencing conformity of the Goods and Related Services	12.3 (c)	Other procurement-specific documents required from Tenderers in ITT Clause 12.3(c) are as follows: Additional Tender Documents are not required.
19	Registration of Goods	11.1(d) &	NONE

	in Tanzania	12.4(a)	
20	Address of Regulators responsible for National Standards and Registration	13.4	**The Director General (DG),** **Tanzania Medicines and Medical Devices Authority(TMDA),** P.O. Box 1253, Dodoma or P.O. Box77150, Dar es Salaam, Tanzania Telephone: +255 22 2450512 /2450751 / 2452108, Fax: +255 22 2450793 Email: info@tmda.go.tz Website: https://www.tmda.go.tz **The Director General (DG),** **Tanzania Bureau of Standards (TBS),** P.O.Box 9524, Sam Nujoma Road / Morogoro Road,Ubungo, Dar es Salaam, Telephone: +255 22 2450206 TollFree Number: 0800110827 Fax: + 25522 2450959, Email: info@tbs.go.tz /Website:https://www.tbs.go.tz/ **The Chief Government Chemist,** **Government Chemist Laboratory Authority,** Physical Address: 5 Barack Obamadrive, Postal Address: P.O. Box 164, Dar es Salaam. Tanzania Tel: +255 22 2113383/4; Fax: +255222113320, Email: gcla@gcla.go.tzWebsite: https://gcla.go.tz/ **The Director General,** **Tanzania Atomic Energy Commission (TAEC),** P.O. Box 743, Block J, Plot No.216, Njiro Arusha, Telephone: +255 272 970050 / 51 /52 / 53 Mobile: +255 754 361221 (DG)

			Fax: +255 272 970054 E-mail : dg@taec.go.tzWebsite: https://www.taec.go.tz
21	Exceptional Tender Document (if any) to be Submitted	13.5(e)	Exceptional documents to be submitted are: NONE
	Lot and Package-Multiple Contracts	14.1	Not Applicable
23	The Lots Coordinator	14.5	Not Applicable
24	Alternative Tenders	16.1	Alternative Tenders to the requirements of the Tendering documents: not allowed
25	Alternative Delivery Schedule	16.2	Alternative Delivery Schedule not allowed.
26	Alternative Specifications	16.3	not allowed
27	Price Adjustment	17.5	NOT_APPLICABLE
28	Tender Prices and Discounts	17.6	Not Applicable
29	Incoterms	17.7	Incoterms 2010
30	Final Destination	17.8 (a) (iii),	ALL COUNCILS IN MBEYA REGION

		(b) (i) &(ii) and c (v)	
31	Fixed Budget	17.9	The available budget is N/A.
32	Tender Currencies	18.1(a) & 18.2	The Tanzanian Shilling
33	Exchange Rate	18.3	The rates of exchange to be used by the Tenderer shall be those established by the Bank of Tanzania twenty-eight (28) days prior to the date specified for opening of tenders.
34	Tender Validity Period	19.1	45
35	Tender Security or Tender Securing Declaration	20.1 & 20.3 (b)	Tender Securing Declaration
36	Format and Signing of Tender	21.2	The written confirmation of authorization to sign on behalf of the Tenderer is: POWER OF ATTORNEY

C. Submission of Tenderers

37	Address for Submission of Samples.	22.5	Not Applicable
38	Deadline for submission of tender.	23.1	Tender Submission Date: 21/07/2026 Tender Submission Time: 10:00 AM hours Local Time.

D. Opening and Evaluation of Tenderers

39.	Conversion to Single Currency	30.2	The rates of exchange to be used by the PE for evaluation and comparison purpose shall be those established by the Bank of Tanzania on the date of opening the tender. Currency for Tender Conversion: The Tanzanian Shilling.
40	Evaluation of Items or Lots	31.2(a)	Procedure for evaluation of items or lots are specified in the Section of Qualification and Evaluation Criteria.
41	Other Evaluation Factors	31.7	All evaluation criteria are found in the Section of Qualification and Evaluation Criteria.
42.	Rate of determining Abnormally low tender	37.4	The rate of determination for abnormally low tenders 15
E. Award of Contract			
43.	Percentage to increase/decrease at Time of Award	41.1	0 percent.
44	Performance Security	43.1 & 43.3	Performance Securing Declaration will be used
45	Environmental and Social Performance Security (ESPS)	43.2	Environmental and Social Performance Security is not applicable.
46.	Maximum Amount of	45.1	Not Applicable percent of the agreed contract price.

	Advance Payment		

F. Review of Procurement Decisions

47	Physical Address for Submission of Complaints/Appeals to PPAA	51.4	Complaints/Appeals to PPAA shall be submitted through NeST at: https://nest.go.tz/module/complaint-management/dashboard. The address for the Appeal to PPAA: The Executive Secretary, Public Procurement Appeals Authority, Ministry of Finance and Planning, Mkandarasi Place, 4th Floor Jakaya Kikwete Road P.O. Box 1385, Dodoma Tanzania Telephone +255 26 2962411 Mobile:+255743505505 Fax + 255 022 2120460 Email: info@ppaa.go.tz or es@ppaa.go.tz Website www.ppaa.go.tz

PHARMACEUTICALS

(Additional Clauses)

C. Preparation of Tenders

TDS Clause Number	**Required Information/Data**	**ITT Number**	**Amendments of, and Supplement to, Clauses in the Instruction to Tenderers**
48	Additional Qualification Criteria	11.1(d)	Documentary evidence of the Tenderer's qualifications to perform the Contract if its Tender is accepted: H as a Good Distribution Practice (GDP) Certificate where appropriate. The Tenderer will submit the following additional information: L ist of pharmaceuticals being manufactured by the Tenderer with product registration/license number and date. A certificate of Pharmaceutical Product as recommended by the WHO for each item offered.
49	Other Specific Procurement Documentation.	12.3 (c)	[Sample clauses] The Goods offered should meet the specified pharmacopeial standards as stated in the Technical Specification. If the Goods delivered are not included in one of the specified pharmacopeias (e.g., the case of a new drug), the Tenderer will provide testing protocols and alternative reference standards.

B. Preparation of Tenders

TDS Clause Number	**Required Information/Data**	**ITT Number**	**Amendments of, and Supplement to, Clauses in the Instruction to Tenderers**
50	Additional Qualification Criteria	11.1(d)	*[Sample Clauses*] *Documentary evidence of the Tenderer's qualifications to perform the Contract if its Tender is accepted:* (a) *is certified by a competent authority in the country of manufacture according to resolution WHA 28 65 (2) of the World Health Organization's Certificate Scheme on the Quality of Pharmaceutical Products Moving in International Commerce.* The Tenderer will submit the following additional information: - *list of vaccines being manufactured by the Tenderer with product registration / license number and date*.
51	Other Specific Procurement Documentation	12.3 (c)	*1.* The Goods to be supplied under the Contract must be licensed both in the country of manufacture and in the United Republic of Tanzania by the time of Contract signing by a recognized NCA. An NCA is an organization that performs all six critical functions for control of biological products as defined by the World Health Organization, namely: licensing based on a published set of requirements; surveillance of vaccine field performance; system of lot release for vaccines; use of laboratory when need; regular inspections for good manufacturing practice and evaluation of clinical performance. The license from the country of manufacture must state that the Tenderer is licensed to manufacture the Goods

			by the NCA in the manufacturing country. Documentary evidence in the form of a certified copy of the license and a copy of the vaccine license/registration that the offered vaccine has been licensed by the NCAs of the manufacturer's country shall accompany the Tender and a copy of the license issued by an NCA in the United Republic of Tanzania must be submitted by Contract signing. If there is no NCA with specific biologics expertise in the United Republic of Tanzania, the Tenderer shall furnish evidence that the Goods meet the qualification criteria in the Technical Specifications. 2. *If the Goods offered do not meet the specified pharmacopoeia standards as stated in the Technical Specification, the Tenderer will provide testing protocols and alternative reference standards.*

SECTION IV: QUALIFICATION AND EVALUATION CRITERIA

QUALIFICATION AND EVALUATION CRITERIA

- **Commercial Evaluation**

1. **Eligibility**

- **Litigation History (SCORE: Comply/Not Comply to specified minimum requirements)**
  Tenderers are required to provide litigation records resulting from contracts completed or ongoing under their execution (In case of Joint Venture, compliance requirements are all Parties Combined – Must Meet requirements and Each Member – Must Meet requirements).

Litigation History Start Year	2020-01-01
Litigation History End Year	2026-06-19

- **Completion Period (SCORE: Comply/Not Comply to specified minimum requirements)**
  Tenderers are required to comply with the completion period as proposed by the procuring entity unless alternative completion period is allowed.

Completion Time (Days)	1095

2. **Standard Tender Forms**

- **Tender Validity Period (SCORE: Comply/Not Comply to specified minimum requirements)**
  Suppliers are required to confirm with the tender validity period specified by the Procuring Entity.

Tender Validity Period (Days)	45

- **Notarized Special Power of Attorney (SCORE: Comply/Not Comply to specified minimum requirements)**
  Tenderers must fill in Standard Power of Attorney as per the required format

and upload into the system.

- **Tender Securing Declaration (SCORE: Comply/Not Comply to specified minimum requirements)**
  The tenderer should submit a tender securing declaration as per instructions.

3. **Financial Situation and Performance**

- **Financial Statement (SCORE: Comply/Not Comply to specified minimum requirements)**
  Audited balance sheets or, if not required by the laws of the Tenderer‘s country, other financial statements acceptable to the PE, for mentioned duration shall be submitted and must demonstrate the current soundness of the Tenderer‘s financial position and indicate its prospective long-term profitability. (In case of Joint Venture, compliance requirements are: Each Member – Must Meet requirements).

Financial Statement Start Date	2023-01-01
Financial Statement End Date	2025-12-31
Minimum Current Ratio [Current Assets(CA)/Current Liabilities(CL)]	1
Minimum Cash Ratio [Cash and Bank(C&B)/Current Liabilities(CL)]	N/A
Minimum Working Capital [Current Assets(CA)-Current Liabilities(CL)]	1
Minimum Gross Profit Margin [Gross Profit(GP)/Total Revenue(TR)*100]	10
Maximum Debt to Equity Ratio [Total Liabilities(TL)/Total Equity(TE)]	1
Minimum Return on Assets [Profit	5

before Tax(PBT)/Total Assets(TA)*100]	

- **Average Annual Turnover (SCORE: Comply/Not Comply to specified minimum requirements)**
  Average Annual Turnover of the mentioned amount, calculated as total certified payments received for contracts in progress and/or completed within the mentioned duration. (In case of Joint Venture, compliance requirements are: All Parties Combined – Must Meet requirements, Each Member – Must Meet percentage requirements and if One Member – Must Meet percentage requirements stated).

Average Annual Turnover Amount in TZS or any other freely convertible currency	650000000
Turnover Start Date	2023-01-01
Turnover End Date	2025-12-31

- **Access to Financial Resources (Sources of Fund) (SCORE: Comply/Not Comply to specified minimum requirements)**
  Tenderers are required to demonstrate details of their sources of finance that show their ability to access adequate finances to meet the cash flow requirements of current and future contracts. (In the case of a Joint Venture, compliance requirements are all Parties Combined – Must Meet requirements).

Average fund amount from all sources (any freely convertible currency proposed by bidder)	650000000

- **Technical Evaluation**

1. **Experience**

- **Current Contract Commitment (SCORE: Comply/Not Comply to specified minimum requirements)**
  Tenderer must demonstrate that it has adequate sources of finance to meet the cash flow requirements on contracts currently in progress and for future contract commitments. (In case of Joint Venture, compliance requirements are: All Parties Combined – Must Meet requirements).

Current Commitment Start Year	2026-08-14
Current Commitment End Year	2029-08-13

- **Specific Experience (SCORE: Comply/Not Comply to specified minimum requirements)**
  Specific and Contract Management Experience: A minimum number of similar contracts based on the physical size, complexity, methods/technology and/or other characteristics described in the PE Requirements on contracts that have been satisfactorily and substantially completed (substantial completion shall be based on 80% or more of completed assignments under the contract) as a prime contractor/supplier/service provider, joint venture member, management contractor/supplier/service provider or sub-contractor/supplier/service provider for mentioned duration. (In case of Joint Venture, compliance requirements are: All Parties – Must Meet requirements). In the case of JVCA, the value of contracts completed by its members shall not be aggregated to determine whether the requirement of the minimum value of a single contract has been met. Instead, each contract performed by each member shall satisfy the minimum value of a single contract as required for single entity. In determining whether the JVCA meets the requirement of total number of contracts, only the number of contracts completed by all members each of value equal or more than the minimum value required shall be aggregated.

- **General Experience (SCORE: Comply/Not Comply to specified minimum requirements)**
  Tenderer should provide details of their previous and on-going contracts to

evidence their general experience in supply of goods.

General experience start date	2020-01-01
General experience end date	2026-06-19
Number of contract	1
Contract value in the specified currency	650000000

- **General Experience in Key Activities (SCORE: Comply/Not Comply to specified minimum requirements)**
  Experience in Key Activities: For any other contracts completed and/or under implementation as prime contractor/supplier/service provider, within the duration and with the minimum experience requested for the key activities. The minimum experience requirement for multiple contracts will be the sum of the minimum requirements for respective individual contracts, unless specified otherwise. (In case of Joint Venture, compliance requirements are: All Parties combined – Must Meet requirements and One Member - Must meet the requirements for the key activities listed and the corresponding minimum requirements).

Employer's Name and Contact	RAS MBEYA
Project Description	PROCUREMENT OF MADICINE
Project Amount	7000000000
Project Start Date	2026-08-14
Project End Date	2029-08-13
Key Activities	SUPPLY OF MEDICINE
Roles in Contract	SUPPLY MEDICINE FOR ALL COUNCILS
Key Personnel	REGIONAL MEDICAL OFFICER

- **Experience in Packaging Distribution and Transportation (SCORE:**

Comply/Not Comply to specified minimum requirements)
The Tenderer should provide proof of experience with and knowledge of modes of packing, distribution, and transportation of pharmaceuticals similar to those subject to Tendering under logistical and climatic conditions similar to the ones in the purchaser's country.

Experience in packaging, distribution and transportation	PACKED IN ORIGINAL MANUFACTURER, SHIPPED BY PRIME VENDOR VEHICLES OR CONTRACTED TRASPORTERS
Number of contracts for packaging, distribution and transportation	1
Minimum value of contracts for packaging, distribution and transportation (TZS)	100000

2. **Technical Specifications**

- **Conformance to Technical Specifications and Standards (SCORE: Comply/Not Comply to specified minimum requirements)**
  Tenderers are required to comply with technical requirements (Service specifications, Technology specifications, Security Specifications, Technical Architecture, Usability, Testing and Quality Assurance, Service Specifications, Conformity to Technical requirements).

3. **Delivery Schedule**

- **Delivery Period (SCORE: Comply/Not Comply to specified minimum requirements)**
  Tenderers are required to comply with delivery period specified by the procuring entity unless alternative delivery schedule has been allowed.

Delivery Period	14

4. **Key Personnel**

- **Key Personnel (SCORE: Comply/Not Comply to specified minimum requirements)**
  Tenderer should provide details of their personnel with adequate qualifications as required by the procuring entity.

Categories of Key Personnel	PHARMACEUTICAL PERSONEL
Education Level	BACHELOR DEGREE AND DIPLOMA
Experience of Key Personnel	2
Number of Required Key Personnel	2

- **Financial Evaluation**

1. **Price Schedule**

- **Priced schedule (SCORE: Comply/Not Comply to specified minimum requirements)**
  The tenderer must quote for each item in the schedule of requirements provided by the procuring entity.

Section IV: Qualification and Evaluation Criteria

This Section contains all the criteria that the PE shall use to evaluate a Tender and qualify the Tenderers. In accordance with **ITT3 (Eligible Tenderers), ITT** 2 9 (Preliminary Evaluation of Tenders), **ITT** 30 (Conversion to Single Currency), **ITT** 31 (Detailed Evaluation and Comparison of Tenders), **ITT** 3 2 (National Preference), and **ITT** 3 3 (Exclusive Preference) no other methods, criteria and factors shall be used. Tenderers shall provide all the information requested in the forms included **in Section V (Tendering Forms**).

Wherever a tenderer is required to state a monetary amount, the stated amount shall be indicated in **Tanzanian shillings (TZS).** However, if a tenderer uses different type of currency, he should indicate the currency to its equivalent in Tanzanian shillings (TZS) using the exchange rate that shall be the selling rates for similar transactions established by the Bank of Tanzania (BoT):

For turnover or financial data required for each year – Exchange rate prevailing on the last day of the respective calendar year (in which the amounts for that year is to be converted) was originally established.
Value of single contract– Exchange rate prevailing on the date of the contract signing

1. Margin of Preference

A) National Preference

i. PE shall grant a margin of preference to tenders offering goods mined, produced or manufactured within the United Republic of Tanzania for the purpose of tender comparison in accordance with the procedures outlined in subsequent paragraphs.

ii. All substantially responsive tenders will be classified in one of the three groups, as follows:

(a) Group A: Tenders offering goods mined, manufactured or produced within the United Republic of Tanzania, for which:

(i) labor, raw materials, and components from the United Republic of Tanzania account for more than thirty (30) per cent of the EXW price of the goods offered; and

(ii) the production facility in which they will be manufactured, assembled or processed has been engaged in manufacturing, assembling or processing such goods at least since the time of Tender submission.

(b) Group B: Other tenders offering goods manufactured within Tanzania, where

the labor, raw materials, and components sourced within the United Republic account for less than thirty percent (30%) of the EXW price of the goods offered**.**

(c) Group C: Tenders offering goods from overseas which are to be directly imported.

iii. EXW prices quoted for tenders in Group A and Group B will include all duties and taxes paid or payable for raw materials or components purchased domestically or imported from overseas.

iv. The price quoted for tenders in group C will include the cost of goods, insurance, and freight (CIF) up to the entry port, border, or any other location, excluding customs duties and other import taxes.

v. All evaluated Tenders in each group will then be compared among themselves to determine the lowest evaluated Tender of each group.

vi. The lowest evaluated Tender of each group will next be compared with the lowest evaluated Tenders of the other groups. If this comparison results in a Tender from Group A or Group B being the lowest, it will be selected for contract award.

vii. Subject to paragraph (vi) above, if the lowest evaluated tender comes from Group B, all Group B tenders will then be compared to the lowest evaluated tender from Group A, with a fifteen percent(15%) addition to the lowest evaluated tender price of the portion of the imported goods in each Group B tender. If, after this adjustment, the Group A tender is the lowest, it will be awarded. If not, the lowest evaluated tender from Group B, as determined after the comparison, will be awarded.

viii. If, as a result of paragraph (vi) comparison, the lowest evaluated Tender is from Group C, all Group C Tenders will then be further compared with the lowest evaluated Tender from Group A and Group B, after adding fifteen (15) percent to the evaluated Tender price of the imported goods offered in each Group C and Group B Tenders.

ix. If the Group A or Group B Tender price in the further comparison is the lowest, it will be selected for award. If not, the lowest evaluated Tender from Group C, as determined from the comparison above, will be selected for award.

B) Margin of Preference to local Suppliers whose investment capital falls under the medium or larger enterprises category as per Small and Medium Enterprises National Development Policy, 2003 (as amended) sub-contracting to local Suppliers whose investment capital falls under the micro or small enterprises category as classified by the Policy in tenders under Exclusive Preference.

Subject to **ITT** 33., a margin of preference shall be granted to the local firm sub-contracting as follows: -

(a) where a local supplier whose investment capital falls under the medium or large enterprises category subcontracts between 20%to 30% of the supply and installation works to a supplier whose investment capital falls under the micro or small enterprises category, the local supplier under the medium or large enterprises category shall be granted a margin of preference of 10% .

(b) where a local supplier under the medium or large enterprises category subcontracts between 10% to 19% of the supply and installation works to a local supplier whose investment capital falls under the micro or small enterprises category, the local supplier under the medium or large enterprises category shall be granted a margin of preference of 8%.

(c) where a local supplier under the medium or large enterprises category subcontracts between 1% to 9% of the supply and installation works to a local supplier whose investment capital falls under the micro or small enterprises category, the local supplier under the medium or large enterprises category shall be granted a margin of preference of 5% .

1. Evaluation (ITT 31)

1.1. ***Evaluation Criteria (ITT 31.6)***

The PE's evaluation of a tender may take into account, in addition to the Tender Price quoted in accordance with ITT Clause 1 7 .8, one or more of the following factors as specified in ITT 31 .2 (a) (b) (c) (d) or (e) and referring to ITT 31 .6**,** using the following criteria and methodologies.

(a) Delivery schedule. (as per INCOTERMs specified in the TDS)

The Goods specified in the List of Goods are required to be delivered within the acceptable time range (after the earliest and before the final date, both dates inclusive) specified in Section VII, Schedule of Requirements. No credit will be given to deliveries before the earliest date, and tenders offering delivery after the final date shall be treated as nonresponsive. Within this acceptable period, an adjustment, as specified in the Delivery Schedule , will be added, for evaluation purposes only, to the tender price of tenders offering deliveries later than the "Earliest Delivery Date" specified in Section VII, Schedule of Requirements.

(b) Deviation in payment schedule. *[insert one of the following]*

(i) Tenderers shall state their tender price for the payment schedule outlined in the SCC. Tenders shall be evaluated on the basis of this base price. Tenderers are, however, permitted to state an alternative payment schedule and indicate the reduction in tender price they wish to offer for such alternative payment schedule. The PE may consider the alternative payment schedule and the reduced tender price offered by the Tenderer selected on the basis of the base price for the payment schedule outlined in the SCC.

or

(ii) The SCC stipulates the payment schedule specified by the PE. If a tender deviates from the schedule and if such deviation is considered acceptable to the PE, the tender will be evaluated by calculating interest earned for any earlier payments involved in the terms outlined in the tender as compared with those stipulated in the SCC, at the rate per annum specified in the Delivery Schedule.

(c)Specific additional criteria

Other specific additional criteria to be considered in the evaluation, and the evaluation method shall be detailed in the Delivery Schedule]

1.2. Multiple Contracts (ITT 3 1.5)

The PE shall award multiple contracts to the Tenderer that offers the lowest evaluated combination of tenders

(one contract per tender) and meets the post-qualification criteria (this Section IV, Sub-Section ITT 37.1 Post Qualification Requirements). The PE shall:

(a) evaluate only lots or contracts that include at least the percentages of items per lot and quantity per item as specified in ITT 17.6

Tender evaluation of such tenders will be carried out as per the following procedures. The average price of an item quoted by substantially responsive Tenderers will be added to the tender price of those who did not quote for that item and the equivalent total cost of the tender so determined will be used for tender comparison, evaluation, and award

(b) take into account:

(i) the lowest-evaluated tender for each lot and

(ii) the price reduction per lot and the methodology for its application as offered by the Tenderer in its tender".

2.Qualification

No.	Subject	Requirement	Single Tender	Joint Venture (existing or intended)			Partnership	Submission Requirements /Source of Document
				All Parties Combined	Each Member	One Member		
1.1. Eligibility								
1.1.1.	Nationality	Nationality in accordance with ITT3.1	Must meet requirement	Must meet requirement	Must meet requirement	N/A	Must meet requirement	Forms ELI – 1.1 and 1.2, with attachments
1.1.2.	Conflict of Interest	No conflicts of interest in accordance with ITT3.8	Must meet requirement	Must meet requirement	Must meet requirement	N/A	Must meet requirement	Form of Tender
1.1.3.	Not declared Ineligible	Not having been declared ineligible as described in	Must meet requirement	Must meet requirement	Must meet requirement	N/A	Must meet requirement	Form of Tender

		ITT3.9						
1.1.4.	Government Owned Entity	Meets conditions of ITT3.10	Must meet requirement	Must meet requirement	Must meet requirement	N/A	N/A	Forms ELI – 1.1 and 1.2, with attachments
1.1.5.	Anti-Bribery Policy	Submission of anti-bribery policy/ code of conduct and Compliance Programme	Must meet requirement	Must meet requirement	Must meet requirement	N/A	Must meet requirement	Form-INTEG
1.1.6.	Power of Attorney	Submission of Power of Attorney	Must meet requirement	Must meet requirement	N/A	N/A	Must meet requirement	
1.1.7.	Tender Security	Submission of Tender Security	Must meet requirement	Must meet requirement	N/A	N/A	Must meet requirement	
	Sub-Criteria for Preference Schemes for Individual Person/Sole							

	Proprietor							
1.1.8.	Citizen Identity Card or National Identification Number (NIN)	Submission of Citizen identity card or NIN	Must meet requirement	N/A	N/A	N/A	N/A	From NIDA
	Sub-Criteria for Preference Schemes for Companies							
1.1.9.	BRELA Consolidated form	Submission Consolidated form	Must meet requirement	N/A	N/A	N/A	N/A	Form from BRELA
1.1.10.	Evidence of Profit Sharing arrangements	Submission of proof of Profit Sharing arrangements	Must meet requirement	N/A	N/A	N/A	N/A	
1.1.11.	Dividend distribution Policies	Submission of Dividend distribution Policies	Must meet requirement	N/A	N/A	N/A	N/A	
1.1.12.	Citizen Identity Card or National Identification Number (NIN)	Submission of Citizen Identity	Must meet requirement	N/A	N/A	N/A	N/A	From NIDA

		Card or NIN						
	Sub-Criteria for Preference Schemes for JVCA							
1.1.13.	Letter of Intent or a Joint Venture Agreement	Submission of Letter of Intent or a Joint Venture Agreement	N/A	Must meet requirement	N/A	N/A	N/A	
1.1.14.	JV Registration Certificate	Submission of Registration Certificate	N/A	Must meet requirement	N/A	N/A	N/A	From CRB
1.1.15.	Citizen Identity Card or National Identification Number (NIN)	Submission of Citizen identity card or NIN	N/A	N/A	Must meet requirement	N/A	N/A	From NIDA
	Sub-Criteria for Partnerships in Preference Schemes Arrangement							
1.1.16.	Extract of Partnership Deed	Submission of Extract of Partnership	N/A	N/A	N/A		Must meet requirement	Form from BRELA

		Deed						
1.1.17.	Citizen identity card or National Identification Number (NIN)	Submission of Citizen identity card or NIN	N/A	N/A	N/A	N/A	Must meet requirement	From NIDA
1.2. Historical Contract Non-Performance								
1.2.1.	History of Non-Performing Contracts	Non-performance of a contract2did not occur as a result of contractor default since 1st January [Insert year].	Must meet requirement1, 2	Must meet requirements	Must meet requirement3	N/A		Form CON-2
1.2.2.	Debarment based on Execution of Tender Securing Declaration by the Authority	Not under debarment based on executi	Must meet requirement	Must meet requirement	Must meet requirement	N/A		Tender Submission Form

		on of a Tender Securing Declaration pursuant to ITT 3.9 (e)						

No.	Subject	Requirement	Single Tenderer	Joint Venture (existing or intended)			Submission Requirements
				All Parties Combined	Each Member	One Member	
1.1.1.	Pending Litigation	Tenderer's financial position and prospective long term profitability sound according to criteria established in 3.1 below and assuming that all pending	Must meet requirement	N/A	Must meet requirement	N/A	Form CON – 2

		litigation will be resolved against the Tenderer					
1.1.2.	Litigation History	No consistent history of court/arbitral award decisions against the Tenderer 4 since 1st January [insert year]	Must meet requirement	N/A	Must meet requirement	N/A	Form CON – 2
1.1.3.	Compliance with Statutory Requirements	No consistent history by the Tenderer 5 of failure to pay taxes and social security Contributions, and no failure to comply with	Must meet requirement	N/A	Must meet requirement	N/A	Form CON – 2

		environmental and health and safety requirements since 1st January [insert year]					
1.2. Financial Situation and Performance							
1.2.1.	Financial Capabilities	(i) The Tenderer shall demonstrate that it has access to, or has available, liquid assets, unencumbered real assets, lines of credit, and other financial means (independent of any contractual advance	Must meet requirement	Must meet Requirement	N/A	N/A	Form FIN – 3.1, with attachments

		payment) sufficient to						

ELIGIBILITYSUB-CRITERIA FOR SUB-CONTRACTING ARRANGEMENTS IN PREFERENCE SCHEMES

No.	**Subject**	**Requirement**	**Foreign Firm Requirement**	**Local Firm Requirement**			**Submission Requirements/Source of Document**
	Local Firm Sub-Contracting Local Firm						
1.	**Letter of Intent to Sub-contract**	Submission of **Letter of Intent to Sub-contract**	N/A	N/A			
2.	**BRELA Consolidated form**	Submission Consolidated form	N/A	N/A			**Form from BRELA**
3.	**Evidence of Profit Sharing arrangements**	Submission of proof of **Profit Sharing arrangements**	N/A	N/A			
4.	**Dividend distribution Policies**	Submission of **Dividend distribution Policies**	N/A	N/A			

5.	**Citizen Identity Card or National Identification Number (NIN)**	Submission of **Citizen identity card or NIN**	N/A	N/A			
	Foreign Firm Sub-Contracting Local Firm						
6.	**Letter of Intent to Sub-contract**	Submission of **Letter of Intent to Sub-contract**	Must meet requirements	N/A	N/A	NA	
7.	**BRELA Consolidated form**	Submission Consolidated form	N/A	Must meet requirements	N/A	N/A	**Form from BRELA**
8.	**Evidence of Profit Sharing arrangements**	Submission of proof of **Profit Sharing arrangements**	N/A	Must meet requirement	N/A	N/A	
9.	**Dividend distribution Policies**	Submission of **Dividend distribution Policies**	N/A	Must meet requirement	N/A	N/A	
10.	**Citizen Identity Card or National**	Submission of **Citizen identity**	Must meet requirement	Must meet requirement	N/A	N/A	

	Identification Number (NIN)	**card or NIN**					

1.1. Key Personnel[15]

Tenderer shall provide details of their personnel with adequate qualifications as required by the procuring entity. The Tenderer shall complete the relevant Forms in Section V, Tendering Forms.

Key Personnel Position	**Education Level**	**Experience of Key Personnel**	**Number of Personnel Required**	**Nationality**[16]	**Name of Company/Firm of Personnel**

1.3. Equipment

The Tenderer shall demonstrate ownership or ability to obtain the required key equipment as listed by the procuring entity

SN	**Type of Key Equipment Required**	**Equipment Capacity**	**Start Year of Equipment Manufacture**	**End Year of Equipment Manufacture**	**Number of Equipment**	**Status of ownership (leased, owned or hired)**

The Tenderer shall provide further details of proposed items of equipment using the relevant Form in Section V, Tendering Forms.

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[1] for tenders set aside for capacity building, the compliance requirements are All Parties Combined – Must Meet requirements.

[2] for tenders set aside for capacity building, the compliance requirements are All Parties Combined – Must Meet requirements

[3] for tenders set aside for capacity building, the compliance requirements are All Parties Combined – Must Meet requirements

[4] for tenders set aside for capacity building, the compliance requirements are All Parties Combined – Must Meet requirements

[5] In the case of local firms with no experience, this criterion shall be waived

[6] for tenders set aside for capacity building, the experience of each member shall be aggregated.

[7] for tenders set aside for capacity building, the experience of each member shall be aggregated

[8] In the case of local firms with no experience, this criterion shall be waived

[10] In the case of local firms with no experience, this criterion shall be waived

[11] for tenders set aside for capacity building, the experience of each member shall be aggregated.

[12] In the case of local firms with no experience, this criterion shall be waived

[13] for tenders set aside for capacity building, the experience of each member shall be aggregated

[14] for tenders set aside for capacity building, the experience of each member shall be aggregated

[15]In case of Joint Venture and Partnership, for tenders set aside for capacity building, the evaluation of personnel requirements must be met by all parties combined.

[16] Tenderer should submit citizen identity card or number issued by National Identification Authority (NIDA) to confirm nationality of Key Personnel.

Section VI: Eligible Countries

All countries are eligible except those subject to the following provisions. A country shall not be eligible if:

1. as a matter of law or official regulation, the Government of Tanzania prohibits commercial relations with that country, provided that the Government of Tanzania is satisfied that such exclusion does not preclude effective competition for the provision of goods or related services required; or
2. by an act of compliance with a decision of the United Nations Security Council taken under Chapter VII of the Charter of the United Nations, the Government of Tanzania prohibits any import of goods from that country or any payments to persons or entities in that country.

PART 2 – PROCURING ENTITY'S REQUIREMENTS

Section VII: Schedule of Requirements

The delivery schedule expressed as weeks/months stipulates hereafter a delivery date which is the date of delivery:-

(i) at EXW premises, or

(ii) to the carrier at the port of shipment when the contract is placed on FOB or CIF terms, or

(iii) to the first carrier when the contract is placed on FCA or CIP terms.

To determine the correct date of delivery hereafter specified, the PE has taken into account the additional time that will be needed for international or national transit to the Project Site or to another commonplace.

List of Commodities

Tender Description: Procurement of medicines from prime vendor for Councils health facilities

Tender No: 78/2025/2026/G/72

LOT NO. 78/2025/2026/G/72

Procurement of medicines from prime vendor for Councils health facilities

GFS Code: 22004102 - Drugs and Medicines

S/N	Description	Unit of measure	Quantity
1	CEPHALEXIN 250MG/5ML ORAL SUSPESNSION, 100MLS	Bottle	6
2	LOPERAMIDE 2MG TABLETS (LOPA)	10TB	1000
3	HEPARIN 1000IU/ML, 5ML INJ	5 Vial	10
4	DEXAMETHASONE 0.1% 5ML EYE DROPS (XSONE)	Bottle	6
5	IBUPROFEN 200MG TABLETS	100TB	1000
6	CLARITHROMYCIN 250MG TABLET	10TB	6
7	CHLORPHENIRAMINE 4MG TABLETS	100TB	6
8	CLONIDINE 300MCG TABLET	112TB	6
9	ETAMSYLATE 500MG TABLET.	18TB	100
10	CEFUROXIME 250MG TABS (THEOROXIME)	10TB	6
11	PIROXICAM 20MG CAPSULE	100TB	1000
12	GLIMEPIRIDE 2+ METFORMIN 500MG TABLETS (ILET B2)	30TB	300
13	METFORMIN 500MG TABLETS	100TB	1000

14	ATORVASTATIN 40MG TABLETS	30TB	6
15	CETIRIZINE TABLET 10MG (ALLERIEVE)	100TB	6
16	DIGOXIN 0.25MG TABLETS	100TB	10
17	ACETYLSALICYLIC ACID (ASPIRIN) 300MG TABLET	100TB	1
18	SALBUTAMOL 100MCG INHALER	1PC	20
19	BORIC ACID EAR DROPS	Bottle	6
20	TAMSULOSIN 0.4MG TABLETS	30TB	30
21	CIPROFLOXACIN 500MG+ TINIDAZOLE 600MG TABLETS	10TB	6
22	PARACETAMOL 1000MG INJECTION (LENIOL)	VIAL	3000
23	CHLORPROMAZINE 100MG TABLETS	100TB	6
24	MAGNESIUM SULPHATE 250MG/ML (50%) 10MLS SOLUTION FOR INJECTION	VIAL	600
25	METFORMIN 500MG + GLIBLENCLAMIDE 5MG TABLET	100TB	1000
26	TELMISARTAN 80MG + HYDROCHLOROTHIAZIDE 12.5MG TABLETS (SAFETELMI-H)	30TB	30
27	ACTINOMYCINE D 0.5 MG INJECTION	AMP	1
28	ASCORBIC ACID 100MG TABLETS	100TB	6
29	NIFEDIPINE 20MG TABLETS	100TB	3000
30	DICLOFENAC 50 MG + PARACETAMOL 500 MG TABLET	100TB	4000
31	PROMETHAZINE HCL 5MG/5ML ORAL LIQUID, 60ML	Bottle	100
32	BETAMETHASONE + SALICYLIC	Tube	6

	OINTMENT, 15GM		
33	TIZANIDINE 4MG TABLETS	10TB	30
34	SODIUM CROMOGLYCATE 2% EYE DROPS, 10MLS	Bottle	300
35	ARTESUNATE 60MG POWDER FOR INJECTION	VIAL	6
36	HYDROCORTISONE 100MG, POWDER FOR INJECTION	VIAL	1000
37	CLOTRIMAZOLE PESSARY WITH APPLICATOR 100 MG	6CP	6
38	SILVER NITRATE STICK	EACH	100
39	BETAMETHASONE 0.025%+CLOTRIMAZOLE 1%+GENTAMICIN0.5% (SKDERM) 30GM	1TU	6
40	HYDROCORTISONE ACETATE EYE DROPS 0.5%	Bottle	1000
41	ARTEMETHER/LUMEFANTRINE 20/120 MG TABLETS	540TB	6
42	FLUCLOXACILLIN 125MG +AMOXICILLIN 125M, 250MG/5ML , 100ML	Bottle	200
43	GLUCOSAMINE SULPHATE 500MG+ CHONDROITIN SULPHATE 400MG TABLETS	30TB	6
44	AZITHROMYCIN 500MG TABLET	3TABS	6
45	BRIMONIDINE EYE DROP 0.2%, 10 MLS (BRIZAGAN)	Bottle	6
46	CANDESARTAN TABLETS 16MG (CANDEZ 16)	30TB	6
47	MUPIROCIN 2% CREAM, 15GM	Tube	3000

48	ANTI - RABIES IMMUNOGLOBULIN 1000IU/5MLS INJECTION	1 Vial	6
49	TRANEXAMIC ACID 500MG TABLET	100TB	300
50	GENTAMYCIN SULPHATE EYE DROPS, 0.3%, 10MLS	Bottle	1000
51	CLOBETASOL PROPIONATE 0.05% OINTMENT, 20GM	Tube	6
52	INSUIN MIXTARD (NOVOMIX) 3ML	VIAL	100
53	OXYBUTYNIN 5MG CAPSULE	P/28	100
54	ESOMEPRAZOLE 40MG TABLETS	14TB	100
55	FLUCLOXACILLIN 250MG+ AMOXYCILLIN 250MG CAPSULES	20CAP	1000
56	AMPICILLIN 250MG+ CLOXACILLIN 250MG CAPSULES	100TB	6
57	AZITHROMYCIN DEHYDRATE 250MG TABLETS	6CP	6
58	INSULIN HUMAN SOLUBLE VIAL 100IU/ML, 10 ML INJECTION	VIAL	100
59	DEXAMETHASONE 4MG TABLETS	100TB	6
60	CO-TRIMOXAZOLE 480MG TABLETS	100TB	6
61	AMOXICILLIN 125MG DISPERSIBLE TABLETS	100TB	6
62	FUSIDIC ACID CREAM 15G	Tube	10
63	FLUCONAZOLE 2MG/ ML,100MLS INJECTION	VIAL	1000
64	LINDANE (1%)+BENZOCAINE (2%) CREAM 25G	Tube	100
65	CEFTRIAXONE 1000 MG +SULBACTAM 500 MG POWDER FOR INJECTION (NEOMICEF)	VIAL	6

66	VITAMIN A - RETINOL CAPSULES 200,000 IU CAPSULES	100TB	20
67	PHENOBARBITAL 30MG TABLETS	100TB	300
68	ALLOPURINOL 300MG TABLET	30CP	6
69	DEXAMETHASONE 4 MG/ML INJECTION	AMP	10
70	CAPTOPRIL 25MG TABLETS	100TB	6
71	VITAMIN A (RETINOL) 100,000 I.U CAPSULES	100TB	50
72	BENZOYL PEROXIDE 5% GEL, 20GM	Tube	6
73	OXYTETRACYCLINE 1% EYE OINTMENT	Tube	3000
74	CLOMIPHENE 50MG TABLET	10TB	6
75	MULTIVITAMIN COMBINATION TABLET	100TB	300
76	SULPHADOXINE 500MG +PYRIMETHAMINE 25MG TABLETS	100TB	30
77	TRAMADOL 50MG CAPSULES	100TB	300
78	CRESOL SAPONATED (LYSOL) 50% LIQUID	Galon	6
79	HYDRALAZINE 20MG/ML 1ML INJECTION	5 Vial	100
80	ACYCLOVIR EYE OINTMENT	1TU	1
81	FEBUXOSTAT 40MG TABLETS	30TB	100
82	TERBINAFINE 15G CREAM 1%	Tube	50
83	LATANOPROST 0.005% EYE SOLUTION	Bottle	30
84	ERYTHROMYCIN 250MG TABLETS	100TB	100
85	METRONIDAZOLE 200MG TABLETS	100TB	3000
86	MEBENDAZOLE SUSPENSION 100MG/5ML	Bottle	800
87	FLUDROCORTISONE 0.1MG TABLETS	10TB	10

88	SODIUM LACTATE COMPOUND,500MLS ,SOLUTION FOR INTRAVENOUS INFUSSION	24BT	300
89	MUPIROCIN 2% OINTMENT 15GM	1TU	3000
90	CHLORHEXIDINE 1.5% + CETRIMIDE 15% LIQUID	Galon	6
91	CLONAZEPAM 2MG TABLET	30TB	6
92	CALAMINE LOTION, 100ML	Bottle	6
93	KETAMINE 10ML 50MG/ML INJECTION.	VIAL	10
94	LINEZOLID 400MG TABLET	30TB	300
95	MICONAZOLE 40GM VAGINAL CREAM (GYNAZOL)	Tube	400
96	BETAMETHASONE (VALERATE) OINTMENT OR CREAM 0.025%	1TU	6
97	BETAMETHASONE + GENTAMYCIN EYE DROPS	Bottle	6
98	ISOFLURANE INHALATION 250ML BOTTLE	Bottle	60
99	FERROUS FUMARATE/ANTI ANAEMIA SYRUPS 20MG/ML, 200 MLS (HEMOVIT)	Bottle	1000
100	FLUPHENAZINE DECANOATE 25 MG/ML INJECTION	10AMP	100
101	CLOTRIMAZOLE 2% VAGINAL CREAM 30GM	Tube	6
102	ENALAPRIL 5MG TABLETS	30TB	10
103	ALBENDAZOLE SUSPENSION 100MG/5ML IN 10ML BOTTLE	Bottle	6
104	CIPROFLOXACIN 200MG/100ML INJECTION	AMP	6

105	METOCLOPRAMIDE 10MG TABLETS	200TB	500
106	TIMOLOL 0.5% EYE DROPS, 5MLS	Bottle	30
107	DANAZOL 100MG TABLET	100TB	6
108	ANTI-HAEMORRHOIDS SUPPOSITORIES	10PC	6
109	BENZATHINE PENICILLIN FORTIFIED PDR F INJ 2.4 MU	50VL	6
110	AMPICILLIN + CLOXACILLIN 125MG/5ML SUSPENSION	24BT	6
111	LIDOCAINE 5%+DEXTROSE 7.5 % SPINAL INJECTION.	10AMP	1000
112	WATER FOR INJECTION 10ML	100PC	30
113	PHYTOMENADIONE(VITAMIN K1) INJECTION	AMP	3000
114	ALUMINIUM HYDROXIDE355MG + MAGNESIUM HYDROXIDE 60MG + SIMETHICONE 100MG + LIQUORICE 400MG (RELCER GEL) 180MLS	Bottle	6
115	SILVER SULFADIAZINE 1% + CHLORHEXIDINE 20GM CREAM	Tube	300
116	HYOSCINE -N- BUTYLBROMIDE 10MG TABS	100TB	100
117	CLOTRIMAZOLE CREAM 1% (PROCOSTAT)	Tube	6
118	HYDROCORTISONE 1%, 15 GRAMS, CREAM	Tube	100
119	ATROPINE 1MG/ML, 1ML, INJECTION.	10AMP	6
120	FLUCYTOSINE 250MG TABLET	10TB	10
121	LIVOLIN FORTE (MULTIVITAMIN) CAPSULE	30TB	300

122	BACLOFEN 10MG TABLETS	30CP	6
123	HALOTHANE INHALATION	Bottle	20
124	HYDROCORTISONE 0.01%,15G CREAM	Tube	1000
125	MIDAZOLAM 5MG/ML INJECTION	AMP	30
126	MICONAZOLE NITRATE 400MG PESSARIES (GYNAZOL)	3TB	3000
127	METFORMIN 850MG TABLET	P/90	300
128	PREGABALIN 75MG CAPSULE	30CP	200
129	LACTULOSE 3.4GM/5ML SOLUTION	Bottle	50
130	CARBAMAZAPINE 200MG TABLETS	100TB	6
131	MOMETASONE NASAL SPRAY	Bottle	1000
132	DOXYLAMINE SUCCINATE 10MG + PYRIDOXINE HCL 10MG TABLETS	10TB	100
133	DIAZEPAM 5MG TABLETS	100TB	6
134	ISOSORBIDE DINITRATE 10MG TABLETS	100TB	100
135	FLUCLOXACILLIN 250MG CAPSULE	10CP	10
136	CLONIDINE 500µG/ML INJECTION.	VIAL	6
137	DICLOFENAC SODIUM 50MG TABLET	100TB	100
138	CLOPIDOGREL 75MG TABLETS	30TB	6
139	BETAMETHASONE 0.1 + NEOMYCIN 0.5% CREAM	Bottle	6
140	RABIES VACCINE USP (POTENCY OF RABIES ANTIGEN ≥ 2.5 IU/DOSE	VIAL	10
141	ENALAPRIL 2.5MG TABLET	30TB	30
142	EPLERENONE 50 MG TABLET	30TB	100
143	BETAMETHASONE 0.1%, 15GM CREAM	Tube	6

144	GENTIAN VIOLET 30ML	Bottle	100
145	CARBAMAZEPINE 100MG TABLET	100TB	6
146	KETOPROFEN 50MG CAPSULES	EACH	1000
147	CALCIUM + VITAMIN D3+VITAMIN A CAPSULE	50PC	6
148	VITAMIN B COMPLEX TABLET.	100TB	20
149	IPRATROPIUM NEBULIZER SOLUTION 500MCG/2ML (IPRAVENT)	Bottle	6
150	VITAMIN B COMPLEX SYRUP, 100ML	24BT	20
151	BUDESONIDE INHALER 100MCG PER DOSE (200 DOSES)	Bottle	6
152	ALUMINIUM HYDROXIDE + MAGNESIUM HYDROXIDE SUSPENSION, 100MLS	24BT	6
153	ORAL REHYDRATION SALTS (ORS) FOR 1 LITRE POWDER	Sachet	1000
154	TELMISARTAN 40MG TABLET	30TB	30
155	MEFENAMIC ACID 500MG TABLET	100TB	300
156	MANNITOL 20% 100MLS SOLUTION FOR IV INFUSION (ABTOL)	Bottle	3000
157	LABETALOL 10MG/ML INJECTION	10VL	30
158	GLIMEPIRIDE 1+ METFORMIN 500MG TABLETS (GEMER 1)	30TB	500
159	PRAZIQUANTEL 600MG TABLET	100TB	200
160	CANDERSARTAN 8MG TABLET	30TB	6
161	METRONIDAZOLE 200MG/5ML 100ML SUSPENSION	24BT	300
162	LORAZEPAM 2MG/ML IN 1ML AMPOULE INJECTION	AMP	50

163	BENZYL PENICILLIN PDR F 5 MU INJECTION	50VL	6
164	CARVEDILOL 6.25MG TABLET	28TB	6
165	ALBENDAZOLE 200MG TABLETS	2TABS	900
166	ARTEMETHER/LUMEFANTRINE 20/120 MG TABLETS	180TABS	6
167	LOW MOLECULAR WEIGHT HEPARIN 25MG INJECTION	AMP	10
168	GLICLAZIDE 40MG TABLET	100TB	100
169	METRONIDAZOLE 1% GEL (CORNIZOLE)	Tube	100
170	HYDROGEN PEROXIDE MOUTHWASH 3% SOLUTION, 100MLS	Bottle	150
171	DEXTROSE 5% (D5) INTRAVENOUS INFUSION 500MLS	24BT	6
172	SALBUTAMOL 4MG TABLETS	100TB	1000
173	ATENOLOL 50MG TABLETS	30TB	6
174	TERBINAFINE 250MG TABLET.	30TB	30
175	BENZOYL PEROXIDE 2.5% ,20GM GEL	1TU	6
176	PROMETHAZINE 25MG TABLETS	100TB	500
177	PARACETAMOL 125MG SUPPOSITORIES	10SUP	300
178	CLARYTHROMYCIN 500MG + TINIDAZOLE 500MG + LANSOPRAZOLE 30MG (42 TABLETS	Kit	6
179	HYDROGEN PEROXIDE 3%, 10MLS EAR DROPS SOLUTION	Bottle	10
180	ESOMEPRAZOLE 20MG TABLET	100TB	3000
181	CEFIXIME 400MG TABLETS	10TB	6
182	MONTELUKAST 10MG TABLETS	30TB	300

183	BISACODYL 5MG TABLET	100TB	6
184	BUPIVACAINE 0.5% (HEAVY MARCAINE) HYPERBARIC, 4ML INJECTION	VIAL	6
185	LEVOTHYROXINE 25MCG TABLET	30TB	30
186	BISOPROLOL FUMARATE 10 MG TABLET	28TB	6
187	CALCIUM GLUCONATE 100MG/ML 10ML INJECTION	10VL	6
188	CEFIXIME 200MG TABLET	10CP	6
189	POTASSIUM PERMANGANATE 1:4000,100ML SOLUTION	Bottle	140
190	CHLORAMPHENICOL 5% EAR/EYE DROPS	Bottle	6
191	BENZOIC ACID COMPOUND (WHITFIELD) OINTMENT 6%+3%	1TU	6
192	GENTAMYCIN 40MG/ML, 2ML INJECTION	10AMP	1000
193	DESLORATIDINE 5MG TABLETS	30TB	6
194	PODOPHYLINE TOPICAL SOLUTION 10-25%	Bottle	20
195	ASCORBIC ACID 500MG TABLET	30TB	6
196	CLOTRIMAZOLE EAR DROPS 15MLS	Bottle	6
197	BRIMONIDINE TARTATE 0.2% EYE DROPS	Bottle	6
198	ECONAZOLE 5% EYE DROP	Bottle	10
199	ACYCLOVIR 10G CREAM	1TU	1
200	CEPHALEXIN 250MG CAPSULES	100TB	6
201	FLUCONAZOLE 150MG TABLETS	100TB	100
202	DIMERCAPSULEROL 50MG/ML 2ML INJECTION	AMP	10
203	PHENYTOIN 100MG TABLETS	100TB	300

204	DIHYDROGESTERONE 10MG TABLETS	10TB	100
205	BUPIVACAINE (LIGHT MARCAINE) 5MG/5ML INJECTION	VIAL	6
206	BUDESONIDE INHALER 200MCG PER DOSE (200 DOSES)	Bottle	6
207	MICONAZOLE CREAM 2%, 20GM	Tube	60
208	OXYTOCIN 10IU/ML, 1ML INJECTION	10AMP	4000
209	ANTI RHO-D HUMAN IMMUNOGLOBIN MONOCLONAL 150MCG INJECTION	AMP	6
210	DONEPEZIL 5MG TABLET	30TB	10
211	FUROSEMIDE 40MG TABLETS	100TB	100
212	CEFTAZIDIME SODIUM 250MG INJECTION	VIAL	6
213	IMIPRAMINE 25MG TABLET	10TB	100
214	CLOPIDOGREL 300MG TABLET	100TB	6
215	ATROPINE 1% EYE DROPS	Bottle	6
216	EPHEDRINE 30MG/ML, 1ML INJECTION	10AMP	1
217	BROMOCRIPTINE MESYLATE 2.5MG TABLET	100TB	6
218	ARTEMETHER/LUMEFANTRINE 20/120 MG TABLETS	720TB	6
219	DEXTROSE 50%, 250ML SOLUTION FOR INTRAVENOUS INJECTION/INFUSION	10PC	6
220	FLUOXETINE 20MG CAPSULE	20CP	10
221	TIMOLOL 0.25% EYE DROPS, 10MLS	Bottle	30
222	DICLOFENAC SODIUM 25MG/ML, 3ML INJECTION	10AMP	10
223	SALBUTAMOL 2.5MG NEBULIZER	Bottle	200

224	GLICLAZIDE 60MG TABLETS	100TB	100
225	ISOSORBIDE MONONITRATE 10MG TABLET	100TB	100
226	PHENOBARBITAL SODIUM 100MG/ML 2ML INJECTION	VIAL	1000
227	CO-PACK FOR DIARRHOEA TREATMENT KIT (ORS+ZINC 20MG) TABLET	Kit	6
228	DICLOFENAC GEL 30G, 1% W/W	Tube	12
229	AMPICILLIN + CLOXACILLIN POWDER FOR INJECTION 500MG	50VL	4
230	GLIBENCLAMIDE 5MG TABLETS	100TB	1000
231	DOBUTAMINE 12.5 MGS/ML INJECTION 20 MLS.	AMP	10
232	HYDROXYCOBALAMINE(VITAMIN B 12) 1000 MCG,INJECTION.	AMP	100
233	SILVER SULPHADIAZINE 1% W/W CREAM	Tube	200
234	CHLORAMPHENICOL + DEXAMETHASONE EYE DROPS, 5MLS	Bottle	6
235	LIGNOCAINE HYDROCHLORIDE INJECTION 1% 10ML	VIAL	100
236	CHLORPHENIRAMINE 10MG/ML INJECTION, 1ML	10AMP	6
237	CHLORPROMAZINE 25MG TABLET	100TB	6
238	BENZOYL PEROXIDE 10% CREAM, 20GM	Tube	6
239	METHYLDOPA 250MG TABLETS	100TB	1000
240	ACTIVATED CHARCOAL 200 MG (WELLVITA)	Tablet	1
241	DEXAMETHASONE 0.1% + CHLORAMPHENICOL EYE DROP 1%W/V	Bottle	6

	5ML		
242	CARBIMAZOLE 5MG TABLETS	100TB	6
243	CETIRIZINE HYDROCHLORIDE 5MG/5ML ORAL LIQUID	Bottle	14
244	2,3-DIMERCAPSULETO-SUCCINIC ACID 100MG CAPSULES	Pack	1
245	LORAZEPAM 2 MG TABLETS (LORIVAN 2)	20TABS	50
246	ONDANSETRON 8MG TABLET	100TB	300
247	DILTIAZEM 60MG TABLETS.	100TB	10
248	SPIRONOLACTONE 25MG TABLETS	100TB	20
249	HYDROGEN PEROXIDE 6% 100ML	Bottle	100
250	FINASTERIDE 5MG TABLETS	20TABS	100
251	TELMISARTAN 80MG TABLET	30TB	30
252	ERYTHROMYCIN 125MG/5ML 100ML	Bottle	100
253	INSULIN HUMAN ZINC SUSPENSION INJECTION 100IU (LENTE)	VIAL	100
254	MEBENDAZOLE 100MG TABLETS	100TB	5000
255	ACYCLOVIR 200MG TABLETS	100TB	20
256	AMOXYCILLIN + CLAVULANIC ACID INJECTION 1.2GM INJECTION	Bottle	6
257	NIFEDIPINE 10MG SLOW RELEASE TABLET	100TB	200
258	DEXAMETHASONE 0.5MG TABLET	100TB	6
259	MANNITOL 20% 500MLS INJECTION	AMP	1
260	CYCLOPENTOLATE 1% EYE DROP	Bottle	6
261	CALCIUM 500MG+ VITAMIN D 10MCG TABLET	30CP	6

262	HEPARIN 5000IU/ML, 5MLS INJECTION	VIAL	10
263	MICONAZOLE 2% ORAL GEL 30G	Dozen	30
264	LIGNOCAINE 2% , 50ML INJECTION	1 Vial	1000
265	LANSOPRAZOLE 30MG TABLET	30TB	500
266	CANDESARTAN 16MG + HYDROCHLOROTHIAZIDE 12.5MG TABLET	30TB	6
267	IPRATROPIUM BROMIDE AEROSOL 20MCG INHALATION (AEROSOL)	Bottle	6
268	BENDROFLUAZIDE 5MG TABLETS	100TB	6
269	COUGH EXPECTORANT CHILD SYRUP100ML	Bottle	24
270	PANTOPRAZOLE 40MG INJECTION (PEPRASOL)	AMP	3000
271	DEXTROSE NORMAL SALINE (DNS) INJECTION 500 MLS	24BT	6
272	AMLODIPINE 5MG TABLETS	30TB	6
273	COUGH EXPECTORANT ADULT SYRUP,100MLS	Bottle	24
274	SODIUM VALPROATE 200MG TABLETS	100TB	20
275	TRAMADOL 50MG/ML, 2ML INJECTION	10AMP	100
276	METOCLOPRAMIDE 5MG/5ML 100ML SYRUP	Bottle	100
277	PANCURONIUM BROMIDE 2MG/ML 2ML INJECTION	10AMP	30
278	PROPRANOLOL 40MG TABLET	100TB	30
279	NORFLOXACIN 400MG TABLET	10TB	3000
280	EPHEDRINE NASAL DROPS 0.5%	P/12	100

	(PAEDIATRIC)		
281	DICYCLOMINE HCL 20MG + PARACETAMOL 500MG TABLETS	10TB	100
282	BENZHEXOL 5MG TABLETS	100TB	6
283	TRIAMCINOLONE INJECTION 40MG/ML (TRICORT 40)	VIAL	20
284	DOXYCYCLINE 100MG CAPSULES	100TB	100
285	SODIUM CHLORIDE 0.9% 500ML SOLUTION FOR INTRAVENOUS INFUSSION, 500ML	24BT	200
286	PHENOBARBITAL 100MG TABLETS	100TB	1000
287	LINDANE LOTION 100ML	Tube	100
288	MONTELUKAST 5MG TABLETS	30TB	300
289	CALCIUM GLUCONATE 500MG TABLET	100PC	6
290	ISOSORBIDE DINITRATE 20MG TABLET	P/50	100
291	CARVEDILOL 12.5MG TABLETS	28TB	6
292	ZINC SULPHATE 20MG TABLETS	100TB	30
293	FERROUS SULPHATE 200MG + FOLIC ACID 0.25 MG TABLETS	100TB	2000
294	HYDRALAZINE 25MG TABLETS	100TB	30
295	CALCIUM CARBONATE 500MG TABLETS	75TB	6
296	CHLORPROMAZINE 25MG/ML 2ML INJECTION	10AMP	6
297	CALCIUM + MAGNESIUM + ZINC + VITAMIN D3 TABLET	30TB	6
298	BISOPROLOL FUMARATE 5MG TABLETS	28TB	6
299	METRONIDAZOLE 5MG/ML, 100ML SOLUTION FOR INTRAVENOUS	10VL	3000

	INJECTION/INFUSION		
300	DICLOFENAC SODIUM 100MG SLOW RELEASE	10TB	10
301	PROMETHAZINE HCL 25MG/ML, 2ML INJECTION	10AMP	40
302	FLUMAZENIL INJ	AMP	10
303	BENZYL BENZOATE EMULSION (BBE) 25%W/V	Bottle	6
304	CYCLOPENTOLATE 0.5% EYE DROP	Bottle	6
305	MELOXICAM 15MG TABLETS	100PC	1000
306	CLOTRIMAZOLE 1%W/W, 30GM POWDER	Bottle	6
307	CLINDAMYCIN IV 150MG/ML INJECTION,2MLS	AMP	6
308	FRUSEMIDE 10MG/ML, 2ML INJECTION	10AMP	1000
309	CLARITHROMYCIN 500MG TABLET	14TB	6
310	PROPOFOL 1% MG/ML INJECTION	AMP	20
311	DORZOLAMIDE 2% EYE DROPS (OCUDOR)	Bottle	10
312	DOXAZOCIN 2MG TABLET	100TB	100
313	ACETAZOLAMIDE 250MG TABLET	100TB	1
314	HALOPERIDOL 1.5MG TABLETS	100TB	100
315	DEXTROSE SOLUTION 10%,500ML INTRAVENOUS INFUSION	24BT	6
316	SODIUM DICHLOROISOCYANURATE 2.7GM TABLETS	100TB	30
317	HYOSCINE BUTYLBROMIDE 20MG/ML, 1ML AMPOULE	AMP	100
318	POVIDONE IODINE LIQUID 10%, 250ML	Bottle	2000

319	ANTITHYMOCITE GLOBULIN 50MG/ML, 5ML (ATG) INJECTION	VIAL	6
320	KETOPROFEN 30GM GEL (2.5% w/w)	Tube	300
321	NITROFURANTOIN 100MG TABLETS	100TB	3000
322	GRISEOFULVIN 500MG TABLETS	100TB	300
323	FLUCONAZOLE 200MG TABLET	100TB	100
324	CIPROFLOXACIN 0.3% EYE DROPS 10MLS	Bottle	6
325	IRBESARTAN 300MG TABLET	30TB	100
326	BETAHISTIDINE 8 MG (BETASERC)TABS	100TB	6
327	GENTAMYCIN 2ML INJ 10MG/ML	10AMP	100
328	GLUTARALDEHYDE WITH ACTIVATOR 2.45% w/v, 5L SOLUTION	Galon	50
329	GLIMEPIRIDE 2MG TABLET	30TB	100
330	LORAZEPAM 1MG TABS	20TABS	40
331	ATROPINE 0.5% EYE DROP	Bottle	6
332	PREDNISOLONE 0.5% EYE DROPS	Bottle	300
333	CARBAMAZEPINE SYRUP 100MG/5ML, 100ML	Bottle	6
334	MULTIVITAMIN SYRUP 100ML	Bottle	300
335	AMOXICILLIN 250MG CAPSULES	100TB	6
336	ERGOCALCIFEROL (VITAMIN D) 1.25MG (50000IU) CAPSULE	10CP	100
337	ESMOLOL 10MG/ML.10ML	AMP	10
338	CLONIDINE 100MCG TABLET	112TB	6
339	SNAKE VENOM POLYVALENT ANTISERUM 10MLS, VIAL.	VIAL	10
340	CHLORPHENIRAMINE MALEATE SYRUP	Bottle	6

	2MG/5ML, 100ML		
341	NORETHISTERONE 5MG TABLET	30TB	300
342	ACETYLSALICYLIC ACID (ENTERIC COATED) 75MG TABLETS	30CP	500
343	PARACETAMOL SYRUP 120MG/5MLS,100MLS	24BT	1000
344	MAGNESIUM TRISILICATE BP COMPOUND TABLETS	100TB	3000
345	LOSARTAN 50MG TABLET	30TB	3000
346	ALLOPURINOL 100MG TABLET	100TB	6
347	AMLODIPINE 10MG TABLET	30TB	6
348	CLONAZEPAM 0.5MG TABLETS	50PC	6
349	CEFIPIME 1GM INJECTION	VIAL	6
350	FOLIC ACID 5MG TABLETS	100TB	1000
351	METOCLOPRAMIDE 5MG/ML ,2ML INJECTION	10AMP	1000
352	TRETINOIN 0.05% CREAM	Tube	5
353	LORATADINE 10MG TABLETS (LORFAST)	30TB	3000
354	LEVODOPA 100 + CARBIDOPA 25MG TABLETS	100TB	50
355	PHENOXYMETHYLPENICILLIN 250MG TABLETS	100TB	1000
356	SUXAMETHONIUM CHLORIDE 50MG/ML INJECTION, 2ML	10AMP	20
357	N-ACETYLCYSTEINE 200MG/ML, 10ML INJECTION	AMP	30
358	BETAMETHASONE 4MG/ML INJECTION	AMP	6
359	PANTOPRAZOLE 40MG TABLETS	30CP	1000

360	CLINDAMYCIN 150MG CAPSULE	100PC	6
361	TELMISARTAN 40 MG + HYDROCHLOROTHIAZIDE 12.5 MG TABLET	30TB	30
362	CHLORAMPHENICOL 1% EYE DROP	Bottle	6
363	CIPROFLOXACIN 500MG TABLETS	100TB	6
364	LORATADINE 5MG/5ML SYRUP (LORATA)	Bottle	2000
365	DOMPERIDONE 10MG TABLETS	10TB	10
366	ADRENALINE 1MG/ML, 1ML INJECTION	10AMP	1
367	IBUPROFEN 100MG/5ML SYRUP 100ML	Bottle	300
368	AMOXICILLIN 500MG + CLAVULANIC ACID 125MG (AUGUMENTIN 625) TABLETS	14TB	1000
369	PREDNISOLONE 5MG TABLETS	100TB	3000
370	AMITRYPTILINE 25MG TABLETS	100TB	6
371	LAMOTRIGINE 100MG TABLETS	60CP	10
372	EPHEDRINE NASAL DROPS 1% (ADULT), 15MLS	P/12	100
373	IBUPROFEN 400MG CAPSULES (IBUBERG)	100TB	100
374	DEXMEDETOMIDINE 200MG/2ML INJECTION	10AMP	6
375	CIPROFLOXACIN EAR DROPS	Bottle	6
376	AMIODARONE 50MG/ML, 3MLS INJECTION	AMP	6
377	Flucytocine 500mg Tablet	100TB	10
378	MEFENAMIC ACID 250MG CAPSULES	100TB	1000
379	DIAZEPAM 5 MG/ML, 2ML INJECTION	10AMP	6

380	CALCIUM +VITAMIN B+ VITAMIN C TABLET	60CP	6
381	TETRACYCLINE EYE OINTMENT 1% 3.5GM	Tube	300
382	CEFTRIAXONE 1G POWDER FOR INJECTION	VIAL	6
383	PHENOXYMETHYLPENICILLIN POWDER FOR ORAL SOLUTION 125MG/5ML	Bottle	2000
384	RABIES VACCINE USP (POTENCY OF RABIES ANTIGEN ≥ 2.5 IU/DOSE	VIAL	30
385	CHLOROXYLENOL (DETTOL) LIQUID 5%	Galon	6
386	BETAXOLOL 0.25% - 0.5% EYE DROP	Bottle	6
387	METOPROL 50 MG TABLETS (METAPROL)	30TB	300
388	FLUTICASONE FUORATE NASAL SPRAY	Bottle	30
389	KETOPROFEN 75MG CAPSULES	30CP	100
390	TACROLIMUS 0.1% OINTMENT	Tube	30
391	CALCIUM CHLORIDE INJECTION 10%, 10MLS	10AMP	6
392	ESOMEPRAZOLE 20MG TABLET	14TB	100
393	LIGNOCAINE HEAVY SPINAL 5%	VIAL	1000
394	XYLOMETAZOLINE 0.1%,10 MLS NASAL SPRAY (XYLOACINO)	Bottle	20
395	ATORVASTATIN 20MG TABLETS	30TB	6
396	ALBENDAZOLE 400MG TABLETS	100TB	2
397	FORMALDEHYDE 36-37% LIQUID	Galon	20
398	ITRACONAZOLE 100 MG CAPSULE	40TABS	50
399	NEOSTIGMINE 2.5MG/ML, 1ML INJECTION	AMP	30

400	LEVETIRACETAM 250MG TABLET	100TB	6
401	AMOXICILLIN + CLAVULIN ACID (228MG/5ML) 100ML, SYRUP	Bottle	6
402	ATROPINE 0.5MG/ML, 1ML, INJECTION.	10AMP	6
403	ERGOCALCIFEROL (VITAMIN D) 0.25MG/ML (10000IU/ML) ORAL SOLUTION	Bottle	10
404	PARACETAMOL 500MG TABLETS	100TB	10000
405	DICYCLOMINE 20MG INJECTION	AMP	100
406	THIOPENTAL SODIUM 500MG, PDR F INJ	25PC	20
407	LISINOPRIL 10MG TABLET	28TB	300
408	FEBUXOSTAT 80MG TABLET	10TB	100
409	NYSTATIN SUSPENSION 100000 IU/ML 30ML	Bottle	3000
410	LOSARTAN 50MG + HYDROCHLORTHIAZIDE 12.5MG TABLETS	30TB	3000
411	ATRACURIUM 50MG/5ML, 5ML INJECTION	AMP	6
412	EPLERENONE 25 MGS, TABLET	30TB	100
413	DICLOFENAC POTASSIUM 50MG TABLET	20TABS	19
414	DEXAMETHASONE 0.1% + NEOMYCIN 0.35% EAR/EYE DROPS (DEXANEO), 10MLS	Bottle	6
415	BETAMETHASONE + GENTAMYCIN EYE DROPS	Bottle	6
416	OLANZAPINE 10MG TABLETS	30TB	300
417	CHLORAMPHENICOL 0.5% EYE OINTMENT	Tube	99
418	CALCIUM +VITAMIN A CAPSULE	10CP	6

419	HEPARIN GEL	Tube	10
420	COAL TAR 5% TOPICAL SOLUTION	Bottle	6
421	CO-TRIMOXAZOLE SUSPENSION 200/40MG/5ML, 100MLS	24BT	6
422	ATORVASTATIN 10MG TABLET	30TB	6
423	POWDER WASHING MACHINE SOAP (SILEX)	Pack	300
424	GLIMEPIRIDE 1MG TABLET	30TB	100
425	IRBESARTAN 150MG TABLETS (IROVEL)	30TB	100
426	DEFEROXAMINE 500MG POWDER FOR (MESYLATE) INJECTION	VIAL	6
427	TRANEXAMIC ACID 100MG/ML 5ML INJECTION	AMP	300
428	ARTEMETHER/LUMEFANTRINE 20/120 MG TABLETS	360TB	6
429	VITAMIN B COMPLEX 2ML INJECTION	10AMP	30
430	AMIODARONE 100MG TABLET	100TB	6
431	GENTAMYCIN 20MG/ML, 2ML INJECTION	10AMP	100
432	ADENOSINE 3MG/ML IN SALINE INJECTION	AMP	1
433	HAND SANITIZER ANTISEPTIC, 250ML	Bottle	2000
434	TINIDAZOLE 500MG TABLET	100TB	100
435	MEROPENEM TRIHYDRATE 1GM INJECTION (POWDER FOR RECONSTITUTION)	VIAL	100
436	AZITHROMYCIN SUSPENSION 200MG/5MLS 15ML	Bottle	6
437	URSODEOXYCHOLIC ACID 300MG	P/50	1

	TABLET		
438	DUTASTERIDE 0.5MG CAPSULE	100TB	10
439	HYDROCHLOROTHIAZIDE 25MG TABLETS	100TB	10

List of Specifications

Tender Description: Procurement of medicines from prime vendor for Councils health facilities

Tender No: 78/2025/2026/G/72

LOT NO. 78/2025/2026/G/72

Procurement of medicines from prime vendor for Councils health facilities

GFS Code: 22004102 - Drugs and Medicines

CEPHALEXIN 250MG/5ML ORAL SUSPESNSION, 100MLS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet and measuring cup. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

LOPERAMIDE 2MG TABLETS (LOPA)

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

HEPARIN 1000IU/ML, 5ML INJ

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. The vials should be packaged separately in individual packs of 5’s. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo to be printed on each vial, and on both, secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

DEXAMETHASONE 0.1% 5ML EYE DROPS (XSONE)

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

IBUPROFEN 200MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CLARITHROMYCIN 250MG TABLET

Technical Requirements	Description

Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CHLORPHENIRAMINE 4MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items

	directly delivered at msd must be registered by TMDA

CLONIDINE 300MCG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ETAMSYLATE 500MG TABLET.

Technical Requirements	Description
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Specification	BP/USP Compendium or any other compendium recognized by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if locally manufactured.
Identification Mark	Labeling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.

CEFUROXIME 250MG TABS (THEOROXIME)

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

PIROXICAM 20MG CAPSULE

Technical Requirements	Description
Specification	BP/USP Compendium. Blister package. Leaflet and Certificate of analysis of the product, TMDA import permit and registration certificate must be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT print on the cap and/or body of the capsule, and GOT and MSD logo on the primary pack and secondary pack
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and

	Mft.dates, Batch No. Weight, size and units per carton.

GLIMEPIRIDE 2+ METFORMIN 500MG TABLETS (ILET B2)

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

METFORMIN 500MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of

	Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ATORVASTATIN 40MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CETIRIZINE TABLET 10MG (ALLERIEVE)

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

DIGOXIN 0.25MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with

	temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ACETYLSALICYLIC ACID (ASPIRIN) 300MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

SALBUTAMOL 100MCG INHALER

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.

Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on bottle and both on secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

BORIC ACID EAR DROPS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

TAMSULOSIN 0.4MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.

Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CIPROFLOXACIN 500MG+ TINIDAZOLE 600MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

PARACETAMOL 1000MG INJECTION (LENIOL)

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CHLORPROMAZINE 100MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.

Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

MAGNESIUM SULPHATE 250MG/ML (50%) 10MLS SOLUTION FOR INJECTION

Technical Requirements	Description
Specification	The product must be registered by TMDA. During tendering must submit the following; -Valid Certificate of medicine registration/Market authorization issued by TMDA - The product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%. Technical documents to be submitted during delivery of consignment: -Valid certificate of analysis -TMDA importation permit for products manufactured outside Tanzania. - Valid Certificate of medicine registration/Market authorization issued by TMDA. -The remained Shelf life of the product during delivery should be not less than 80% of the total shelf life for items with total shelf life of 24 Months or less. For product with total shelf life of more than 24 Months (e.g 36 Months, 48 Months or 60 Months) the delivery shelf life should not be less than 24 Months).
Identification Mark	Labelling must comply with TMDA requirements and include the GOT and MSD logo on all packages (Primary, secondary and tertiary packages)
Shipping Carton	-Five ply carton with MSD and GOT logo.

	-The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. -Supplier vehicle for items directly delivered at MSD must be registered by TMDA.

METFORMIN 500MG + GLIBLENCLAMIDE 5MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

TELMISARTAN 80MG + HYDROCHLOROTHIAZIDE 12.5MG TABLETS (SAFETELMI-H)

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.

Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ACTINOMYCINE D 0.5 MG INJECTION

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ASCORBIC ACID 100MG TABLETS

Technical Requirements	Description

Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

NIFEDIPINE 20MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.

Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

DICLOFENAC 50 MG + PARACETAMOL 500 MG TABLET

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Blister package, leaflet, Certificate of analysis,Import Permit for overseas product or registration permit for local product to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

PROMETHAZINE HCL 5MG/5ML ORAL LIQUID, 60ML

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet and measuring cup. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT

	and MSD logo on bottle and both on secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

BETAMETHASONE + SALICYLIC OINTMENT, 15GM

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet. Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tube and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

TIZANIDINE 4MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.

Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

SODIUM CROMOGLYCATE 2% EYE DROPS, 10MLS

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ARTESUNATE 60MG POWDER FOR INJECTION

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium

	recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

HYDROCORTISONE 100MG, POWDER FOR INJECTION

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items

	directly delivered at msd must be registered by TMDA

CLOTRIMAZOLE PESSARY WITH APPLICATOR 100 MG

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

SILVER NITRATE STICK

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.

Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

BETAMETHASONE 0.025%+CLOTRIMAZOLE 1%+GENTAMICIN0.5% (SKDERM) 30GM

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

HYDROCORTISONE ACETATE EYE DROPS 0.5%

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ARTEMETHER/LUMEFANTRINE 20/120 MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items

	directly delivered at msd must be registered by TMDA

FLUCLOXACILLIN 125MG +AMOXICILLIN 125M, 250MG/5ML , 100ML

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet and measuring cup. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on bottle and both on secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

GLUCOSAMINE SULPHATE 500MG+ CHONDROITIN SULPHATE 400MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity

	75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

AZITHROMYCIN 500MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

BRIMONIDINE EYE DROP 0.2%, 10 MLS (BRIZAGAN)

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must

	be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CANDESARTAN TABLETS 16MG (CANDEZ 16)

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

MUPIROCIN 2% CREAM, 15GM

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet.Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ANTI - RABIES IMMUNOGLOBULIN 1000IU/5MLS INJECTION

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit for imported product and copy of TMDA registration certificate must be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo to be printed on each vial, and on both, secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton

	must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

TRANEXAMIC ACID 500MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

GENTAMYCIN SULPHATE EYE DROPS, 0.3%, 10MLS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CLOBETASOL PROPIONATE 0.05% OINTMENT, 20GM

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA

INSUIN MIXTARD (NOVOMIX) 3ML

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. During transport storage temperature must be as per manufacturer recommendations.
Shipping carton	Five ply carton with MSD and GOT logo. The carton

specifications	must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

OXYBUTYNIN 5MG CAPSULE

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit for imported product and copy of TMDA registration certificate must be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT print on the cap and/or body of the capsule, and GOT and MSD logo on the primary, Secondary and Tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ESOMEPRAZOLE 40MG TABLETS

Technical Requirements	Description
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported

	and/or certificate of registration if locally manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.

FLUCLOXACILLIN 250MG+ AMOXYCILLIN 250MG CAPSULES

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

AMPICILLIN 250MG+ CLOXACILLIN 250MG CAPSULES

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of

	Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

AZITHROMYCIN DEHYDRATE 250MG TABLETS

Technical Requirements	Description
Technical Specifications	BP/USP Compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Package	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

INSULIN HUMAN SOLUBLE VIAL 100IU/ML, 10 ML INJECTION

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium

	recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

DEXAMETHASONE 4MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity,

	UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CO-TRIMOXAZOLE 480MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

AMOXICILLIN 125MG DISPERSIBLE TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of

	Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

FUSIDIC ACID CREAM 15G

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet. Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tube and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

FLUCONAZOLE 2MG/ ML,100MLS INJECTION

Technical Requirements	Description

Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

LINDANE (1%)+BENZOCAINE (2%) CREAM 25G

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet. Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tube and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CEFTRIAXONE 1000 MG +SULBACTAM 500 MG POWDER FOR INJECTION (NEOMICEF)

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. The vials should be packaged separately in individual packs of 10's. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo to be printed on each vial, and on both, secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

VITAMIN A - RETINOL CAPSULES 200,000 IU CAPSULES

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity,

	UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

PHENOBARBITAL 30MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ALLOPURINOL 300MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported

	and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

DEXAMETHASONE 4 MG/ML INJECTION

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CAPTOPRIL 25MG TABLETS

Technical Requirements	Description
Identification Mark on the	Labelling must comply with TMDA requirements.

product	GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

VITAMIN A (RETINOL) 100,000 I.U CAPSULES

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity,

	UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

BENZOYL PEROXIDE 5% GEL, 20GM

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet. Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tube and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

OXYTETRACYCLINE 1% EYE OINTMENT

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet. Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tube and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity,

	UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CLOMIPHENE 50MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

MULTIVITAMIN COMBINATION TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

SULPHADOXINE 500MG +PYRIMETHAMINE 25MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

TRAMADOL 50MG CAPSULES

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity

	75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CRESOL SAPONATED (LYSOL) 50% LIQUID

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Certificate of analysis of the product to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on both packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

HYDRALAZINE 20MG/ML 1ML INJECTION

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not

	exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ACYCLOVIR EYE OINTMENT

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet. Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tube and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

FEBUXOSTAT 40MG TABLETS

Technical Requirements	Description
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted.

	Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.

TERBINAFINE 15G CREAM 1%

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tube and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

LATANOPROST 0.005% EYE SOLUTION

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton

	must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ERYTHROMYCIN 250MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

METRONIDAZOLE 200MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate

	of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

MEBENDAZOLE SUSPENSION 100MG/5ML

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet and measuring cup. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet and measuring cup. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.

Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

FLUDROCORTISONE 0.1MG TABLETS

Technical Requirements	Description
Specification	The product must be registered by TMDA. During tendering must submit the following; -Valid Certificate of medicine registration/Market authorization issued by TMDA - The product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%. Technical documents to be submitted during delivery of consignment: -Valid certificate of analysis -TMDA importation permit for products manufactured outside Tanzania. - Valid Certificate of medicine registration/Market authorization issued by TMDA. -The remained Shelf life of the product during delivery should be not less than 80% of the total shelf life for items with total shelf life of 24 Months or less. For product with total shelf life of more than 24 Months (e.g 36 Months, 48 Months or 60 Months) the delivery shelf

	life should not be less than 24 Months).
Identification Mark	Labelling must comply with TMDA requirements and include the GOT and MSD logo on all packages (Primary, secondary and tertiary packages)
Shipping Carton	-Five ply carton with MSD and GOT logo. -The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. -Supplier vehicle for items directly delivered at MSD must be registered by TMDA.

SODIUM LACTATE COMPOUND,500MLS ,SOLUTION FOR INTRAVENOUS INFUSSION

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Each Polypropylene bottle with an infusion site should be packed in its secondary pack. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items

	directly delivered at msd must be registered by TMDA

MUPIROCIN 2% OINTMENT 15GM

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet.Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CHLORHEXIDINE 1.5% + CETRIMIDE 15% LIQUID

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Certificate of analysis of the product to be submitted.
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo on bottle and both on secondary packages
Shipping carton	Five ply carton with MSD and GOT logo. The carton

specifications	must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CLONAZEPAM 2MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CALAMINE LOTION, 100ML

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.

Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

KETAMINE 10ML 50MG/ML INJECTION.

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

LINEZOLID 400MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA

	import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

MICONAZOLE 40GM VAGINAL CREAM (GYNAZOL)

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

	Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

BETAMETHASONE (VALERATE) OINTMENT OR CREAM 0.025%

Technical Requirements	Description
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet.Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.

BETAMETHASONE + GENTAMYCIN EYE DROPS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT

	and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ISOFLURANE INHALATION 250ML BOTTLE

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

FERROUS FUMARATE/ANTI ANAEMIA SYRUPS 20MG/ML, 200 MLS (HEMOVIT)

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium

	recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

FLUPHENAZINE DECANOATE 25 MG/ML INJECTION

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and

	Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CLOTRIMAZOLE 2% VAGINAL CREAM 30GM

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tube and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ENALAPRIL 5MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with

	temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ALBENDAZOLE SUSPENSION 100MG/5ML IN 10ML BOTTLE

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with leaflet,measuring cups, Certificate of analysis,Import Permit for overseas product or registration Permit for local product to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on bottle and both on secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CIPROFLOXACIN 200MG/100ML INJECTION

Technical Requirements	Description
Technical Specification	BP/USP compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on vial and both on primary and secondary packages
Package	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton

METOCLOPRAMIDE 10MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity

	75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

TIMOLOL 0.5% EYE DROPS, 5MLS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

DANAZOL 100MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on

	primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ANTI-HAEMORRHOIDS SUPPOSITORIES

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognized by TMDA. Leaflet and Certificate of analysis to be submited. Product should have TMDA import permit if imported and/or Registration Certificate if locally manufactured
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

BENZATHINE PENICILLIN FORTIFIED PDR F INJ 2.4 MU

Technical Requirements	Description
Specification	BP/USP compendium or any other Compendium recognised by TMDA. The vial should be packaged separately in individual packs of 50’s. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Print GOT and MSD logo on each vial , and both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton

	must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

AMPICILLIN + CLOXACILLIN 125MG/5ML SUSPENSION

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet and measuring cup. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

LIDOCAINE 5%+DEXTROSE 7.5 % SPINAL INJECTION.

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium

	recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

WATER FOR INJECTION 10ML

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on vial and both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

PHYTOMENADIONE(VITAMIN K1) INJECTION

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA

	import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ALUMINIUM HYDROXIDE355MG + MAGNESIUM HYDROXIDE 60MG + SIMETHICONE 100MG + LIQUORICE 400MG (RELCER GEL) 180MLS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and

	Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

SILVER SULFADIAZINE 1% + CHLORHEXIDINE 20GM CREAM

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet. Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tube and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

HYOSCINE -N- BUTYLBROMIDE 10MG TABS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity

	75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CLOTRIMAZOLE CREAM 1% (PROCOSTAT)

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tube and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

HYDROCORTISONE 1%, 15 GRAMS, CREAM

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet.Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be

	submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ATROPINE 1MG/ML, 1ML, INJECTION.

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

FLUCYTOSINE 250MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

LIVOLIN FORTE (MULTIVITAMIN) CAPSULE

Technical Requirements	Description
Specification	BP/USP Compendium. Blister package. Leaflet and Certificate of analysis of the product, TMDA import permit and registration certificate must be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT print on the cap and/or body of the capsule, and GOT and MSD logo on the primary pack and secondary pack
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and

	Mft.dates, Batch No. Weight, size and units per carton.

BACLOFEN 10MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

HALOTHANE INHALATION

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.

Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

HYDROCORTISONE 0.01%,15G CREAM

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet. Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tube and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

MIDAZOLAM 5MG/ML INJECTION

Technical Requirements	Description
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on vial and both on primary and secondary packages
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on vial and both on primary and secondary packages
Specification	BP/USP compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Specification	BP/USP compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.

MICONAZOLE NITRATE 400MG PESSARIES (GYNAZOL)

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and

	tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

METFORMIN 850MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

PREGABALIN 75MG CAPSULE

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of

	Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

LACTULOSE 3.4GM/5ML SOLUTION

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet and measuring cup. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items

	directly delivered at msd must be registered by TMDA

CARBAMAZAPINE 200MG TABLETS

Technical Requirements	Description
Technical Specifications	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Packaging	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

MOMETASONE NASAL SPRAY

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet. Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tube and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

DOXYLAMINE SUCCINATE 10MG + PYRIDOXINE HCL 10MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

DIAZEPAM 5MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements.

	GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ISOSORBIDE DINITRATE 10MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

FLUCLOXACILLIN 250MG CAPSULE

Technical Requirements	Description

Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Specification	BP/USP Compendium or any compandium recognized by TMDA . Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT print on the cap and/or body of the capsule, and GOT and MSD logo on the primary pack and secondary pack

CLONIDINE 500µG/ML INJECTION.

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

DICLOFENAC SODIUM 50MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CLOPIDOGREL 75MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

BETAMETHASONE 0.1 + NEOMYCIN 0.5% CREAM

Technical Requirements	Description

Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet. Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tube and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

RABIES VACCINE USP (POTENCY OF RABIES ANTIGEN ≥ 2.5 IU/DOSE

Technical Requirements	Description
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA
Specification	BP/USP Compendium or any other recognised compendium. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.

ENALAPRIL 2.5MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

EPLERENONE 50 MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

BETAMETHASONE 0.1%, 15GM CREAM

Technical Requirements	Description

Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet. Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tube and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

GENTIAN VIOLET 30ML

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CARBAMAZEPINE 100MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of

	Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

KETOPROFEN 50MG CAPSULES

Technical Requirements	Description
Specification	BP/USP Compendium. Blister package. Leaflet and Certificate of analysis of the product, TMDA import permit and registration certificate must be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT print on the cap and/or body of the capsule, and GOT and MSD logo on the primary pack and secondary pack
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CALCIUM + VITAMIN D3+VITAMIN A CAPSULE

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet,

	Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

VITAMIN B COMPLEX TABLET.

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

IPRATROPIUM NEBULIZER SOLUTION 500MCG/2ML (IPRAVENT)

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine

	registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on bottle and both on secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

VITAMIN B COMPLEX SYRUP, 100ML

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. TEach bottle should be packed in its secondary pack with a measuring cup and a a leaflet. Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

BUDESONIDE INHALER 100MCG PER DOSE (200 DOSES)

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.

Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on bottle and both on secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ALUMINIUM HYDROXIDE + MAGNESIUM HYDROXIDE SUSPENSION, 100MLS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet and measuring cup. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ORAL REHYDRATION SALTS (ORS) FOR 1 LITRE POWDER

Technical Requirements	Description

Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	Glucose Anhydrous13.5g + Sodium Chloride 2.6g + Potassium Chloride 1.5g + Sodium Citrate2.9g. BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

TELMISARTAN 40MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and

	Mft.dates, Batch No. Weight, size and units per carton.

MEFENAMIC ACID 500MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

MANNITOL 20% 100MLS SOLUTION FOR IV INFUSION (ABTOL)

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and

	Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

LABETALOL 10MG/ML INJECTION

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. The vials should be packaged separately in individual packs of 10's. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo to be printed on each vial, and on both, secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

GLIMEPIRIDE 1+ METFORMIN 500MG TABLETS (GEMER 1)

Technical Requirements	Description
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate

	of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.

PRAZIQUANTEL 600MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CANDERSARTAN 8MG TABLET

Technical Requirements	Description
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

METRONIDAZOLE 200MG/5ML 100ML SUSPENSION

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. The product must be supplied with a measuring cup. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

LORAZEPAM 2MG/ML IN 1ML AMPOULE INJECTION

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on vial and both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

BENZYL PENICILLIN PDR F 5 MU INJECTION

Technical Requirements	Description

Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CARVEDILOL 6.25MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ALBENDAZOLE 200MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ARTEMETHER/LUMEFANTRINE 20/120 MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with

	temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

LOW MOLECULAR WEIGHT HEPARIN 25MG INJECTION

Technical Requirements	Description
Specification	The product must be registered by TMDA/TBS. During tendering must submit the following; -Valid Certificate of product registration/Market authorization issued by TMDA/TBS - The product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%. Technical documents to be submitted during delivery of consignment: -Valid certificate of analysis -TMDA/TBS importation permit for products manufactured outside Tanzania. - Valid Certificate of medicine registration/Market authorization issued by TMDA/TBS. -The remained Shelf life of the product during delivery should be not less than 80% of the total shelf life for items with total shelf life of 24 Months or less. For product with total shelf life of more than 24 Months (e.g 36 Months, 48 Months or 60 Months) the delivery shelf life should not be less than 24 Months).
Identification Mark	Labelling must comply with TMDA requirements and include the GOT and MSD logo on all packages (Primary, secondary and tertiary packages)

Shipping Carton	-Five ply carton with MSD and GOT logo. -The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. -Supplier vehicle for items directly delivered at MSD must be registered by TMDA.

GLICLAZIDE 40MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

METRONIDAZOLE 1% GEL (CORNIZOLE)

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.

Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

HYDROGEN PEROXIDE MOUTHWASH 3% SOLUTION, 100MLS

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate of product to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on both packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

DEXTROSE 5% (D5) INTRAVENOUS INFUSION 500MLS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Each Polypropylene bottle with an infusion site should be packed in its secondary pack. Leaflet and Certificate of analysis of the product to be

	submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

SALBUTAMOL 4MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items

	directly delivered at msd must be registered by TMDA

ATENOLOL 50MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

TERBINAFINE 250MG TABLET.

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.

Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

BENZOYL PEROXIDE 2.5% ,20GM GEL

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet. Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tube and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

PROMETHAZINE 25MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity

	75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

PARACETAMOL 125MG SUPPOSITORIES

Technical Requirements	Description
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CLARYTHROMYCIN 500MG + TINIDAZOLE 500MG + LANSOPRAZOLE 30MG (42 TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity,

	UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

HYDROGEN PEROXIDE 3%, 10MLS EAR DROPS SOLUTION

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ESOMEPRAZOLE 20MG TABLET

Technical Requirements	Description
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

Specification	BP/USP Compendium or any other compendium recognized by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if locally manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT print on the cap and/or body of the capsule, and GOT and MSD logo on the primary pack and secondary pack

CEFIXIME 400MG TABLETS

Technical Requirements	Description
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

MONTELUKAST 10MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity,

	UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

BISACODYL 5MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

BUPIVACAINE 0.5% (HEAVY MARCAINE) HYPERBARIC, 4ML INJECTION

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not

	exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

LEVOTHYROXINE 25MCG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

BISOPROLOL FUMARATE 10 MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not

	exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CALCIUM GLUCONATE 100MG/ML 10ML INJECTION

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CEFIXIME 200MG TABLET

Technical Requirements	Description
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

POTASSIUM PERMANGANATE 1:4000,100ML SOLUTION

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CHLORAMPHENICOL 5% EAR/EYE DROPS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA

	registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

BENZOIC ACID COMPOUND (WHITFIELD) OINTMENT 6%+3%

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tube and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

GENTAMYCIN 40MG/ML, 2ML INJECTION

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of

	analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

DESLORATIDINE 5MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

PODOPHYLINE TOPICAL SOLUTION 10-25%

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. TMDA import permit,copy of

	registration certificate, leaflet and Certificate of analysis of the product to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on bottle and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ASCORBIC ACID 500MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CLOTRIMAZOLE EAR DROPS 15MLS

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.

Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on bottle and both on secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

BRIMONIDINE TARTATE 0.2% EYE DROPS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ECONAZOLE 5% EYE DROP

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton

	must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ACYCLOVIR 10G CREAM

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet. Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tube and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CEPHALEXIN 250MG CAPSULES

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.

Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

FLUCONAZOLE 150MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

DIMERCAPSULEROL 50MG/ML 2ML INJECTION

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import

	permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

PHENYTOIN 100MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

DIHYDROGESTERONE 10MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

BUPIVACAINE (LIGHT MARCAINE) 5MG/5ML INJECTION

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Specification	BP/USP compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.

Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on vial and both on primary and secondary packages
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on vial and both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

BUDESONIDE INHALER 200MCG PER DOSE (200 DOSES)

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on bottle and both on secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

MICONAZOLE CREAM 2%, 20GM

Technical Requirements	Description

Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tube and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

OXYTOCIN 10IU/ML, 1ML INJECTION

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ANTI RHO-D HUMAN IMMUNOGLOBIN MONOCLONAL 150MCG INJECTION

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

DONEPEZIL 5MG TABLET

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with

	temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

FUROSEMIDE 40MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CEFTAZIDIME SODIUM 250MG INJECTION

Technical Requirements	Description

Specification	BP/USP compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on vial and both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

IMIPRAMINE 25MG TABLET

Technical Requirements	Description
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.

CLOPIDOGREL 300MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import

	permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ATROPINE 1% EYE DROPS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

EPHEDRINE 30MG/ML, 1ML INJECTION

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import

	permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

BROMOCRIPTINE MESYLATE 2.5MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ARTEMETHER/LUMEFANTRINE 20/120 MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.

Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

DEXTROSE 50%, 250ML SOLUTION FOR INTRAVENOUS INJECTION/INFUSION

Technical Requirements	Description
Specification	BP/USP compendium. Should be packed in plastic collapsible bottle. TMDA import permit for imported product and copy of TMDA registration certificate for product, Certificate of analysis to be submitted.
Identification Mark	Labelling should comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

FLUOXETINE 20MG CAPSULE

Technical Requirements	Description

Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.

TIMOLOL 0.25% EYE DROPS, 10MLS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

DICLOFENAC SODIUM 25MG/ML, 3ML INJECTION

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

SALBUTAMOL 2.5MG NEBULIZER

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on bottle and both on secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity,

	UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

GLICLAZIDE 60MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ISOSORBIDE MONONITRATE 10MG TABLET

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA

	import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

PHENOBARBITAL SODIUM 100MG/ML 2ML INJECTION

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CO-PACK FOR DIARRHOEA TREATMENT KIT (ORS+ZINC 20MG) TABLET

Technical Requirements	Description
Specification	International Pharmacopoiea. Should be registered by TMDA. Leaflet and Certificate of analysis to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

DICLOFENAC GEL 30G, 1% W/W

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet.Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items

	directly delivered at msd must be registered by TMDA

AMPICILLIN + CLOXACILLIN POWDER FOR INJECTION 500MG

Technical Requirements	Description
Specification	BP/USP compendium or any other Compendium recognised by TMDA. The vial should be packaged separately in individual packs of 50's. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Print GOT and MSD logo on each vial , and both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

GLIBENCLAMIDE 5MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity,

	UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

DOBUTAMINE 12.5 MGS/ML INJECTION 20 MLS.

Technical Requirements	Description
Specification	BP/USP compendium or any other Compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

HYDROXYCOBALAMINE(VITAMIN B 12) 1000 MCG,INJECTION.

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit for imported product and copy of TMDA registration certificate must be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo to be printed on each vial, and on both, secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity,

	UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

SILVER SULPHADIAZINE 1% W/W CREAM

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet.Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CHLORAMPHENICOL + DEXAMETHASONE EYE DROPS, 5MLS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary

	pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

LIGNOCAINE HYDROCHLORIDE INJECTION 1% 10ML

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. The vials should be packaged separately in individual packs of 10’s. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo to be printed on each vial, and on both, secondary and tertiary packages
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo to be printed on each vial, and on both, secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Specification	BP/USP compendium or any other compendium recognised by TMDA. The vials should be packaged separately in individual packs of 10’s. Leaflet and Certificate of analysis of the product to be submitted.

	Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.

CHLORPHENIRAMINE 10MG/ML INJECTION, 1ML

Technical Requirements	Description
Specification	BP/USP compendium or any other Compendium recognised by TMDA. The ampoules should be packaged separately in individual packs of 10's. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Specification	BP/USP compendium or any other Compendium recognised by TMDA. The ampoules should be packaged separately in individual packs of 10's. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on both on primary and secondary packages
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and

	Mft.dates, Batch No. Weight, size and units per carton.

CHLORPROMAZINE 25MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

BENZOYL PEROXIDE 10% CREAM, 20GM

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet.Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tube and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity,

	UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA

METHYLDOPA 250MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ACTIVATED CHARCOAL 200 MG (WELLVITA)

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine

	registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

DEXAMETHASONE 0.1% + CHLORAMPHENICOL EYE DROP 1%W/V 5ML

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Each bottle with a cap should be packed in its secondary pack. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on bottle and both on secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

	Supplier vehicle for items directly delivered at msd must be registered by TMDA

CARBIMAZOLE 5MG TABLETS

Technical Requirements	Description
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CETIRIZINE HYDROCHLORIDE 5MG/5ML ORAL LIQUID

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet and measuring cup. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet and measuring cup. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on bottle and both on secondary packages
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on bottle and both on secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton

	must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

2,3-DIMERCAPSULETO-SUCCINIC ACID 100MG CAPSULES

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tube and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

LORAZEPAM 2 MG TABLETS (LORIVAN 2)

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on

	primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ONDANSETRON 8MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

DILTIAZEM 60MG TABLETS.

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.

Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

SPIRONOLACTONE 25MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and

	Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

HYDROGEN PEROXIDE 6% 100ML

Technical Requirements	Description
Specification	BP/USP compendium or any other Compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit must be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on both packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

FINASTERIDE 5MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity,

	UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

TELMISARTAN 80MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ERYTHROMYCIN 125MG/5ML 100ML

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Each bottle with plastic screw cap with dosing cup should be packed in its secondary pack. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted.

	Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

INSULIN HUMAN ZINC SUSPENSION INJECTION 100IU (LENTE)

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on vial and both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

MEBENDAZOLE 100MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate

	of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ACYCLOVIR 200MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd

	must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

AMOXYCILLIN + CLAVULANIC ACID INJECTION 1.2GM INJECTION

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

NIFEDIPINE 10MG SLOW RELEASE TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity

	75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

DEXAMETHASONE 0.5MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

MANNITOL 20% 500MLS INJECTION

Technical Requirements	Description
Specification	BP/USP compendium or any other Compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or

	certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CYCLOPENTOLATE 1% EYE DROP

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CALCIUM 500MG+ VITAMIN D 10MCG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on

	primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

HEPARIN 5000IU/ML, 5MLS INJECTION

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. The vials should be packaged separately in individual packs of 5’s. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo to be printed on each vial, and on both, secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

MICONAZOLE 2% ORAL GEL 30G

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet.Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding

	30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

LIGNOCAINE 2% , 50ML INJECTION

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

LANSOPRAZOLE 30MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CANDESARTAN 16MG + HYDROCHLOROTHIAZIDE 12.5MG TABLET

Technical Requirements	Description
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

Specification	BP/USP Compendium or any other compendium recognized by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if locally manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.

IPRATROPIUM BROMIDE AEROSOL 20MCG INHALATION (AEROSOL)

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on bottle and both on secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

BENDROFLUAZIDE 5MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate

	of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

COUGH EXPECTORANT CHILD SYRUP100ML

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Each bottle with plastic screw cap with dosing cup should be packed in its secondary pack. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

	Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

PANTOPRAZOLE 40MG INJECTION (PEPRASOL)

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

DEXTROSE NORMAL SALINE (DNS) INJECTION 500 MLS

Technical Requirements	Description
Specification	BP/USP compendium. Should be packed in plastic collapsible bottle. TMDA import permit for imported product and copy of TMDA registration certificate for product, Certificate of analysis to be submitted.
Identification Mark	Labelling should comply with TMDA requirements. GOT and MSD logo print on primary pack and

	secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

AMLODIPINE 5MG TABLETS

Technical Requirements	Description
Specification	BP/USP compendium or any other Compendium recognised by TMDA. Leaflet ,TMDA Imported permit for imported items and TMDA registration for local items.Certificate of analysis of the product to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tablet and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

COUGH EXPECTORANT ADULT SYRUP,100MLS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Each bottle with plastic screw cap with dosing cup should be packed in its secondary pack. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted.

	Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

SODIUM VALPROATE 200MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

TRAMADOL 50MG/ML, 2ML INJECTION

Technical Requirements	Description
Specification	BP/USP compendium or any other Compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit must be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

METOCLOPRAMIDE 5MG/5ML 100ML SYRUP

Technical Requirements	Description
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on bottle and both on secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet amd measuring cup. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.

PANCURONIUM BROMIDE 2MG/ML 2ML INJECTION

Technical Requirements	Description
Identification Mark on the	Labelling must comply with TMDA requirements.

product	GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

PROPRANOLOL 40MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

NORFLOXACIN 400MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

EPHEDRINE NASAL DROPS 0.5% (PAEDIATRIC)

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a droper and leaflet. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on bottle and both on secondary packages

Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

DICYCLOMINE HCL 20MG + PARACETAMOL 500MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

BENZHEXOL 5MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised

	compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

TRIAMCINOLONE INJECTION 40MG/ML (TRICORT 40)

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd

	must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

DOXYCYCLINE 100MG CAPSULES

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

SODIUM CHLORIDE 0.9% 500ML SOLUTION FOR INTRAVENOUS INFUSSION, 500ML

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Each Polypropylene bottle with

	an infusion site should be packed in its secondary pack. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

PHENOBARBITAL 100MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

	Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

LINDANE LOTION 100ML

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

MONTELUKAST 5MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CALCIUM GLUCONATE 500MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ISOSORBIDE DINITRATE 20MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CARVEDILOL 12.5MG TABLETS

Technical Requirements	Description

Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ZINC SULPHATE 20MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.

Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

FERROUS SULPHATE 200MG + FOLIC ACID 0.25 MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

HYDRALAZINE 25MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and

	tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CALCIUM CARBONATE 500MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity,

	UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CHLORPROMAZINE 25MG/ML 2ML INJECTION

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CALCIUM + MAGNESIUM + ZINC + VITAMIN D3 TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.

Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

BISOPROLOL FUMARATE 5MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

METRONIDAZOLE 5MG/ML, 100ML SOLUTION FOR INTRAVENOUS INJECTION/INFUSION

Technical Requirements	Description
Identification Mark on the	Labelling must comply with TMDA requirements.

product	GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

DICLOFENAC SODIUM 100MG SLOW RELEASE

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

PROMETHAZINE HCL 25MG/ML, 2ML INJECTION

Technical Requirements	Description
Specification	BP/USP compendium or any other Compendium recognised by TMDA. The ampoules should be packaged separately in individual packs of 10's, Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

FLUMAZENIL INJ

Technical Requirements	Description
Specification	BP/USP compendium or any other Compendium recognised by TMDA. The ampoules should be packaged separately in individual packs of 25's Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

BENZYL BENZOATE EMULSION (BBE) 25%W/V

Technical Requirements	Description
cream	benzyl benzoate

CYCLOPENTOLATE 0.5% EYE DROP

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

MELOXICAM 15MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

	Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.

CLOTRIMAZOLE 1%W/W, 30GM POWDER

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on bottle and both on secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CLINDAMYCIN IV 150MG/ML INJECTION,2MLS

Technical Requirements	Description
Specification	BP/USP compendium or any other Compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on both on primary and secondary packages

Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

FRUSEMIDE 10MG/ML, 2ML INJECTION

Technical Requirements	Description
Specification	The product must be registered by TMDA. During tendering must submit the following; -Valid Certificate of medicine registration/Market authorization issued by TMDA - The product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%. Technical documents to be submitted during delivery of consignment: -Valid certificate of analysis -TMDA importation permit for products manufactured outside Tanzania. - Valid Certificate of medicine registration/Market authorization issued by TMDA. -The remained Shelf life of the product during delivery should be not less than 80% of the total shelf life for items with total shelf life of 24 Months or less. For product with total shelf life of more than 24 Months (e.g 36 Months, 48 Months or 60 Months) the delivery shelf life should not be less than 24 Months).
Identification Mark	Labelling must comply with TMDA requirements and include the GOT and MSD logo on all packages (Primary, secondary and tertiary packages)
Shipping Carton	-Five ply carton with MSD and GOT logo. -The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name,

	Exp.and Mft.dates, Batch No. Weight, size and units per carton. -Supplier vehicle for items directly delivered at MSD must be registered by TMDA.

CLARITHROMYCIN 500MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

PROPOFOL 1% MG/ML INJECTION

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit for imported product and copy of TMDA registration certificate must be submitted.
Specification	BP/USP compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit for imported product and copy of TMDA registration certificate must be submitted.

Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo to be printed on each vial, and on both, secondary and tertiary packages
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo to be printed on each vial, and on both, secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

DORZOLAMIDE 2% EYE DROPS (OCUDOR)

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd

	must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

DOXAZOCIN 2MG TABLET

Technical Requirements	Description
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ACETAZOLAMIDE 250MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

HALOPERIDOL 1.5MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA

	import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

DEXTROSE SOLUTION 10%,500ML INTRAVENOUS INFUSION

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Each Polypropylene bottle with an infusion site should be packed in its secondary pack. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd

	must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

SODIUM DICHLOROISOCYANURATE 2.7GM TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Sodium Dichloroisocyanurate Tab has a strength of 2.7g per tab with available Chlorine equivalent to 1.5g per tab. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

HYOSCINE BUTYLBROMIDE 20MG/ML, 1ML AMPOULE

Technical Requirements	Description
Specification	BP/USP compendium or any other Compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit must be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT

	and MSD logo on both on primary, secondary and and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

POVIDONE IODINE LIQUID 10%, 250ML

Technical Requirements	Description
Specification	BP/USP compendium. Certificate of analysis, leaflet, TMDA import permit, copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling should comply with TMDA requirements. GOT and MSD logo print on both packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ANTITHYMOCITE GLOBULIN 50MG/ML, 5ML (ATG) INJECTION

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.

Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

KETOPROFEN 30GM GEL (2.5% w/w)

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet. Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tube and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

NITROFURANTOIN 100MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine

	registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

GRISEOFULVIN 500MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

FLUCONAZOLE 200MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CIPROFLOXACIN 0.3% EYE DROPS 10MLS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

IRBESARTAN 300MG TABLET

Technical Requirements	Description

Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.

BETAHISTIDINE 8 MG (BETASERC)TABS

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

GENTAMYCIN 2ML INJ 10MG/ML

Technical Requirements	Description
Specification	BP/USP compendium or any other Compendium recognised by TMDA. The ampoules should be packaged separately in individual packs of 10's. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

GLUTARALDEHYDE WITH ACTIVATOR 2.45% w/v, 5L SOLUTION

Technical Requirements	Description
Identification Mark	Labelling must comply with TMDA requirements and include the GOT and MSD logo on all packages (Primary, secondary and tertiary packages)
Specification	The product must be registered by TMDA. During tendering must submit the following; -Valid Certificate of medicine registration/Market authorization issued by TMDA - The product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%. Technical documents to be submitted during delivery of consignment: -Valid certificate of analysis

	-TMDA importation permit for products manufactured outside Tanzania. - Valid Certificate of medicine registration/Market authorization issued by TMDA. -The remained Shelf life of the product during delivery should be not less than 80% of the total shelf life for items with total shelf life of 24 Months or less. For product with total shelf life of more than 24 Months (e.g 36 Months, 48 Months or 60 Months) the delivery shelf life should not be less than 24 Months).
Shipping Carton	-Five ply carton with MSD and GOT logo. -The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. -Supplier vehicle for items directly delivered at MSD must be registered by TMDA.

GLIMEPIRIDE 2MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

LORAZEPAM 1MG TABS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ATROPINE 0.5% EYE DROP

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and

	Mft.dates, Batch No. Weight, size and units per carton.

PREDNISOLONE 0.5% EYE DROPS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CARBAMAZEPINE SYRUP 100MG/5ML, 100ML

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Each bottle with plastic screw cap with dosing cup should be packed in its secondary pack. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity,

	UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA

MULTIVITAMIN SYRUP 100ML

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. The product must be supplied with a measuring cup. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

AMOXICILLIN 250MG CAPSULES

Technical Requirements	Description
Specification	BP/USP Compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.

Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ERGOCALCIFEROL (VITAMIN D) 1.25MG (50000IU) CAPSULE

Technical Requirements	Description
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Specification	BP/USP Compendium. Blister package. Leaflet and Certificate of analysis of the product, TMDA import permit and registration certificate must be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT print on the cap and/or body of the capsule, and GOT and MSD logo on the primary pack and secondary pack

ESMOLOL 10MG/ML.10ML

Technical Requirements	Description
Specification	BP/USP compendium or any other Compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity,

	UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CLONIDINE 100MCG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

SNAKE VENOM POLYVALENT ANTISERUM 10MLS, VIAL.

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on vial and both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CHLORPHENIRAMINE MALEATE SYRUP 2MG/5ML, 100ML

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet and measuring cup. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet and measuring cup. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on bottle and both on secondary packages
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on bottle and both on secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

NORETHISTERONE 5MG TABLET

Technical Requirements	Description

Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ACETYLSALICYLIC ACID (ENTERIC COATED) 75MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

PARACETAMOL SYRUP 120MG/5MLS,100MLS

Technical Requirements	Description
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

MAGNESIUM TRISILICATE BP COMPOUND TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

LOSARTAN 50MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and

	Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ALLOPURINOL 100MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

AMLODIPINE 10MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported

	and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CLONAZEPAM 0.5MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CEFIPIME 1GM INJECTION

Technical Requirements	Description

Specification	BP/USP compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on vial and both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

FOLIC ACID 5MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

METOCLOPRAMIDE 5MG/ML ,2ML INJECTION

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet amd measuring cup. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on bottle and both on secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

TRETINOIN 0.05% CREAM

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and

	Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA

LORATADINE 10MG TABLETS (LORFAST)

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

LEVODOPA 100 + CARBIDOPA 25MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

PHENOXYMETHYLPENICILLIN 250MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with

	temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

SUXAMETHONIUM CHLORIDE 50MG/ML INJECTION, 2ML

Technical Requirements	Description
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo to be printed on each vial, and on both, secondary and tertiary packages
Specification	BP/USP compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit for imported product and copy of TMDA registration certificate must be submitted.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

N-ACETYLCYSTEINE 200MG/ML, 10ML INJECTION

Technical Requirements	Description
Specification	BP/USP compendium or any other Compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT

	and MSD logo on both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

BETAMETHASONE 4MG/ML INJECTION

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit for imported product and copy of TMDA registration certificate must be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo to be printed on each vial, and on both, secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

PANTOPRAZOLE 40MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Specification	BP/USP Compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.

Identification Mark	Labelling must comply with TMDA requirements. GOT print on the cap and/or body of the capsule, and GOT and MSD logo on the primary pack and secondary pack
Identification Mark	Labelling must comply with TMDA requirements. GOT print on the cap and/or body of the capsule, and GOT and MSD logo on the primary pack and secondary pack
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CLINDAMYCIN 150MG CAPSULE

Technical Requirements	Description
Specification	BP/USP Compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT print on the cap and/or body of the capsule, and GOT and MSD logo on the primary, Secondary and Tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

TELMISARTAN 40 MG + HYDROCHLOROTHIAZIDE 12.5 MG TABLET

Technical Requirements	Description

Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CHLORAMPHENICOL 1% EYE DROP

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CIPROFLOXACIN 500MG TABLETS

Technical Requirements	Description
Technical Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or

	certificate of medicine registration if localy manufactured.
identification mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Package	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

LORATADINE 5MG/5ML SYRUP (LORATA)

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet and measuring cup. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

DOMPERIDONE 10MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ADRENALINE 1MG/ML, 1ML INJECTION

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.

Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

IBUPROFEN 100MG/5ML SYRUP 100ML

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet and measuring cup. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

AMOXICILLIN 500MG + CLAVULANIC ACID 125MG (AUGUMENTIN 625) TABLETS

Technical Requirements	Description

Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

PREDNISOLONE 5MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton	Five ply carton with MSD and GOT logo. The carton

specifications	must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

AMITRYPTILINE 25MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

LAMOTRIGINE 100MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.

Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

EPHEDRINE NASAL DROPS 1% (ADULT), 15MLS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and

	Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

IBUPROFEN 400MG CAPSULES (IBUBERG)

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

DEXMEDETOMIDINE 200MG/2ML INJECTION

Technical Requirements	Description
Specification	BP/USP compendium or any other Compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit must be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT

	and MSD logo on both on primary, secondary and and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CIPROFLOXACIN EAR DROPS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

AMIODARONE 50MG/ML, 3MLS INJECTION

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and

	tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA

Flucytocine 500mg Tablet

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

MEFENAMIC ACID 250MG CAPSULES

Technical Requirements	Description
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.

DIAZEPAM 5 MG/ML, 2ML INJECTION

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CALCIUM +VITAMIN B+ VITAMIN C TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

TETRACYCLINE EYE OINTMENT 1% 3.5GM

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet. Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tube and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CEFTRIAXONE 1G POWDER FOR INJECTION

Technical Requirements	Description

Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

PHENOXYMETHYLPENICILLIN POWDER FOR ORAL SOLUTION 125MG/5ML

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.

Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

RABIES VACCINE USP (POTENCY OF RABIES ANTIGEN ≥ 2.5 IU/DOSE

Technical Requirements	Description
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA
Specification	BP/USP Compendium or any other recognised compendium. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.

CHLOROXYLENOL (DETTOL) LIQUID 5%

Technical Requirements	Description
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on both packages

Specification	BP/USP compendium or any other compendium recognised by TMDA. Certificate of analysis of the product to be submitted.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

BETAXOLOL 0.25% - 0.5% EYE DROP

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

METOPROL 50 MG TABLETS (METAPROL)

Technical Requirements	Description
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and

	tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.

FLUTICASONE FUORATE NASAL SPRAY

Technical Requirements	Description
Specification	Leaflet and Certificate of analysis of the product to be submitted. GCLA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by GCLA Supplier vehicle for items directly delivered at msd must be registered by GCLA

KETOPROFEN 75MG CAPSULES

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised

	compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

TACROLIMUS 0.1% OINTMENT

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet.Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton

	must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CALCIUM CHLORIDE INJECTION 10%, 10MLS

Technical Requirements	Description
Specification	BP/USP compendium or any other Compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit must be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on both on primary, secondary and and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ESOMEPRAZOLE 20MG TABLET

Technical Requirements	Description
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Specification	BP/USP Compendium or any other compendium recognized by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if locally manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT

	print on the cap and/or body of the capsule, and GOT and MSD logo on the primary pack and secondary pack

LIGNOCAINE HEAVY SPINAL 5%

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Specification	BP/USP compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on vial and both on primary and secondary packages
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on vial and both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

XYLOMETAZOLINE 0.1%,10 MLS NASAL SPRAY (XYLOACINO)

Technical Requirements	Description

Specification	BP/USP Compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ATORVASTATIN 20MG TABLETS

Technical Requirements	Description
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of

	analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.

ALBENDAZOLE 400MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with leaflet,measuring cups, Certificate of analysis,Import Permit for overseas product or registration Permit for local product to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on bottle and both on secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and

	Mft.dates, Batch No. Weight, size and units per carton.

FORMALDEHYDE 36-37% LIQUID

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Certificate of analysis of the product, TMDA import permit and copy of product registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on both packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ITRACONAZOLE 100 MG CAPSULE

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and

	Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

NEOSTIGMINE 2.5MG/ML, 1ML INJECTION

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

LEVETIRACETAM 250MG TABLET

Technical Requirements	Description
Specification	The product must be registered by TMDA. During tendering must submit the following; -Valid Certificate of medicine registration/Market authorization issued by TMDA - The product must have storage condition stability for climatic zone IV B with temperature not exceeding

	30°C and Relative humidity 75+_5%. Technical documents to be submitted during delivery of consignment: -Valid certificate of analysis -TMDA importation permit for products manufactured outside Tanzania. - Valid Certificate of medicine registration/Market authorization issued by TMDA. -The remained Shelf life of the product during delivery should be not less than 80% of the total shelf life for items with total shelf life of 24 Months or less. For product with total shelf life of more than 24 Months (e.g 36 Months, 48 Months or 60 Months) the delivery shelf life should not be less than 24 Months).
Identification Mark	Labelling must comply with TMDA requirements and include the GOT and MSD logo on all packages (Primary, secondary and tertiary packages)
Shipping Carton	-Five ply carton with MSD and GOT logo. -The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. -Supplier vehicle for items directly delivered at MSD must be registered by TMDA.

AMOXICILLIN + CLAVULIN ACID (228MG/5ML) 100ML, SYRUP

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.

Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on bottle and both on secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ATROPINE 0.5MG/ML, 1ML, INJECTION.

Technical Requirements	Description
Specification	BP/USP compendium or any other Compendium recognised by TMDA. The ampoules should be packaged separately in individual packs of 10's. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ERGOCALCIFEROL (VITAMIN D) 0.25MG/ML (10000IU/ML) ORAL SOLUTION

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. The product must be supplied with a measuring cup. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.

Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

PARACETAMOL 500MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

DICYCLOMINE 20MG INJECTION

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium

	recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

THIOPENTAL SODIUM 500MG, PDR F INJ

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. The vial should be packaged separately in individual packs of 25’s. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Specification	BP/USP compendium or any other compendium recognised by TMDA. The vial should be packaged separately in individual packs of 25’s. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.

Identification Mark	Print GOT and MSD logo on vial and both on primary and secondary packages
Identification Mark	Print GOT and MSD logo on vial and both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

LISINOPRIL 10MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

FEBUXOSTAT 80MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised

	compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

NYSTATIN SUSPENSION 100000 IU/ML 30ML

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity,

	UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

LOSARTAN 50MG + HYDROCHLORTHIAZIDE 12.5MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ATRACURIUM 50MG/5ML, 5ML INJECTION

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.

Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

EPLERENONE 25 MGS, TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

DICLOFENAC POTASSIUM 50MG TABLET

Technical Requirements	Description

Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

DEXAMETHASONE 0.1% + NEOMYCIN 0.35% EAR/EYE DROPS (DEXANEO), 10MLS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

BETAMETHASONE + GENTAMYCIN EYE DROPS

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit and TMDA

	registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary pack and secondary pack.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

OLANZAPINE 10MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CHLORAMPHENICOL 0.5% EYE OINTMENT

Technical Requirements	Description

Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tube and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

CALCIUM +VITAMIN A CAPSULE

Technical Requirements	Description
Specification	BP/USP Compendium. Blister package. Leaflet and Certificate of analysis of the product, TMDA import permit and registration certificate must be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT print on the cap and/or body of the capsule, and GOT and MSD logo on the primary pack and secondary pack
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

HEPARIN GEL

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each tube should be packed in its secondary pack with a leaflet. Certificate of analysis of the product to be submitted. Should have TMDA

	import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on tube and both on secondary and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

COAL TAR 5% TOPICAL SOLUTION

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

CO-TRIMOXAZOLE SUSPENSION 200/40MG/5ML, 100MLS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Each bottle with plastic screw cap with dosing cup should be packed in its secondary pack. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton specifications	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

ATORVASTATIN 10MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton

	must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

POWDER WASHING MACHINE SOAP (SILEX)

Technical Requirements	Description
Specification	Powder soap suitable for machine wash. Should contain non-ionic surfactant, ant-staining agent and de-mineralised water, non- pungent fragrance,oils and any non- toxic agent.
Identification Mark	Labelling must comply with TBS requirements. GOT and MSD logo on both packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

GLIMEPIRIDE 1MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

IRBESARTAN 150MG TABLETS (IROVEL)

Technical Requirements	Description
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.

DEFEROXAMINE 500MG POWDER FOR (MESYLATE) INJECTION

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for 2°C to 8°C. Data logger with USB port for temperature records during transportation must be submitted during delivery of the consignment.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd

	must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

TRANEXAMIC ACID 100MG/ML 5ML INJECTION

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on vial and both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ARTEMETHER/LUMEFANTRINE 20/120 MG TABLETS

Technical Requirements	Description
Identification Mark on the product	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Shipping carton	Five ply carton with MSD and GOT logo. The carton

specifications	must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

VITAMIN B COMPLEX 2ML INJECTION

Technical Requirements	Description
Specification	BP/USP compendium or any other Compendium recognised by TMDA. Leaflet, Certificate of analysis of the product, TMDA import permit must be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on both on primary, secondary and and tertiary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

AMIODARONE 100MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and

	Mft.dates, Batch No. Weight, size and units per carton.

GENTAMYCIN 20MG/ML, 2ML INJECTION

Technical Requirements	Description
Specification	BP/USP compendium or any other Compendium recognised by TMDA. The ampoules should be packaged separately in individual packs of 10's. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

ADENOSINE 3MG/ML IN SALINE INJECTION

Technical Requirements	Description
Specification	BP/USP compendium or any other Compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

HAND SANITIZER ANTISEPTIC, 250ML

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

TINIDAZOLE 500MG TABLET

Technical Requirements	Description
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity,

	UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

MEROPENEM TRIHYDRATE 1GM INJECTION (POWDER FOR RECONSTITUTION)

Technical Requirements	Description
Specification	BP/USP compendium or any other Compendium recognised by TMDA. The ampoules should be packaged separately in individual packs of 10's. Leaflet and Certificate of analysis of the product to be submitted. Product should have TMDA import permit if imported and/or certificate of registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on both on primary and secondary packages
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

AZITHROMYCIN SUSPENSION 200MG/5MLS 15ML

Technical Requirements	Description
Specification	BP/USP compendium or any other compendium recognised by TMDA. Each bottle should be packed in its secondary pack with a leaflet nd measuring cup. Certificate of analysis of the product to be submitted. Should have TMDA import permit if imported and/or certificate of medicine registration if localy manufactured.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo on bottle and both on secondary packages

Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.

URSODEOXYCHOLIC ACID 300MG TABLET

Technical Requirements	Description
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton.
Specification	BP/USP Compendium or any other compendium recognised by TMDA. Blister package. Leaflet, Certificate of analysis of the product, TMDA import permit and copy of TMDA registration certificate to be submitted.
Identification Mark	Labelling must comply with TMDA requirements. GOT embossment on tablet, and GOT and MSD logo print on primary, secondary and tertiary packages.

DUTASTERIDE 0.5MG CAPSULE

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and

	tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

HYDROCHLOROTHIAZIDE 25MG TABLETS

Technical Requirements	Description
Specification	BP/USP Compendium or any other recognised compendium. Blister package. Leaflet and Certificate of analysis of the product to be submitted. TMDA import permit for products manufactured out of Tanzania must be submitted. Certificate of medicine registration must be submitted. Product must have storage condition stability for climatic zone IV B with temperature not exceeding 30°C and Relative humidity 75+_5%.
Identification Mark	Labelling must comply with TMDA requirements. GOT and MSD logo print on primary, secondary and tertiary packages.
Shipping Carton	Five ply carton with MSD and GOT logo. The carton must be well labelled with Product name, Quantity, UOM and its strength. Manufacturer name, Exp.and Mft.dates, Batch No. Weight, size and units per carton. Supplier vehicle for items directly delivered at msd must be registered by TMDA Supplier vehicle for items directly delivered at msd must be registered by TMDA

Delivery Schedule

Tender Description: Procurement of medicines from prime vendor for Councils health facilities

Tender No: 78/2025/2026/G/72

Specific item(s) delivery schedule will be specified on the issuance of the call-off order.

Inspection and Tests

Tender Description: Procurement of medicines from prime vendor for Councils health facilities

Tender No: 78/2025/2026/G/72

Inspection and Tests for specific item(s) will be known on the issuance of the call-off order.

PART 3 – CONDITIONS OF CONTRACT AND CONTRACT FORMS

SECTION VIII: GENERAL CONDITIONS OF CONTRACT (GCC)

GENERAL CONDITIONS OF CONTRACT

These General Conditions of Contract (GCC), read in conjunction with the Special Conditions of Contract (SCC).

S/N	GCC Clause	GCC Clause Number	GCC Clause Description
1.	Definitions	1.1	In this Contract, the following terms shall be interpreted as indicated: a) "**Arbitrator**" is the person appointed by the appointing authority specified in the **SCC**, to resolve contractual disputes. b) **"Commencement Date"** is the date when the supplier shall commence execution of the contract as specified in Certificate of Contract Commencement issued by the Contract Manager. c) **"Completion Date"** means the date of fulfilment of the related services by the Supplier in accordance with the terms and conditions set forth in the contract and certified by the Project Manger in accordance with GCC 21.1. d) **"Contract"** means an agreement entered between the Purchaser and the Supplier to execute the Contract. e) **"Contract Effectiveness Date"** means the date shown in the Certificate notice of Contract Commencement issued by the Purchaser Contract Manager upon fulfilment of the conditions precedent stipulated in GCC 3. f) **"Contract Price"** is the amount accepted in the Letter of Acceptance and thereafter as adjusted in accordance with the provisions of the Contract. g) "**Day**" means calendar day. h) "**Delivery**" means the transfer of the goods from the supplier equipment, machinery, and /or other materials which the Supplier is required to supply to the Purchaser

			under Contract. i) "**Effective Date**" means the date on which this Contract becomes effective. j) "**Eligible Country**" means the countries and territories eligible for participation in procurements financed by the specified institution. k) "**End User**" means the organization(s) where the goods will be used, as named in the **SCC**. l) **"Final Destination"** where applicable means the place or places named in the SCC. m) **"Force Majeure"** means an unforeseeable event which is beyond reasonable control of either Party and which makes a Party's performance of its obligations under the Contract impossible or so impractical as to be considered impossible under the circumstances. For the purposes of this Contract, "Force Majeure" means an event which is beyond the reasonable control of a Party, is not foreseeable, is unavoidable, and its origin is not due to negligence or lack of care on the part of a Party, and which makes a Party's performance of its obligations hereunder impossible or so impractical as reasonably to be considered impossible in the circumstances; and includes, but is not limited to, war, riots, civil disorder, earthquake, fire, explosion, storm, flood, epidemics, or other adverse weather conditions, strikes, lockouts or other industrial action (except where such strikes, lockouts or other industrial action are within the power of the Party invoking Force Majeure to prevent), confiscation. n) "**GCC**" means the General Conditions of Contract contained in this section. o) "**Goods**" means all of the pharmaceuticals including nutritional supplement and oral and injectable forms of contraception, vaccines, and condoms that the Supplier is required to supply to the Purchaser under the Contract. p) "**Origin**" means the place where the Goods were

			mined, grown, or produced or from which the Services are supplied. Goods are produced when, through manufacturing, processing, or substantial and major assembly of components, a commercially recognized new produce results that is substantially different in basic characteristics or in purpose or utility from its components. q) "**Project Manager**" is the person (or any other competent person appointed by the Employer and notified to the Contractor, to act in replacement of the Project Manager) who is responsible for supervising the execution of the Works and administering the Contract. Details of the project manager will be available in the Contract Finalization Information Section in the Contract Agreement. r) "**Purchaser**" means the organization purchasing the Goods, as named in the **SCC**. s) "**Registration Certificate**" means the certificate of registration or other documents in lieu thereof establishing that the Goods supplied under the Contract are registered for use in the United Republic of Tanzania in accordance with the Applicable Law. t) "**Related Services**" means those services ancillary to the supply of the Goods, such as transportation and insurance, and any other incidental services, such as installation, commissioning, provision of technical assistance, training, initial maintenance and other such obligations of the Supplier covered under the Contract. u) "**SCC**" means the Special Conditions of Contract. v) "**Services**" means those services ancillary to the supply of the Goods, such as transportation and insurance, and any other incidental services, such as provision of technical assistance, training, and other such obligations of the Supplier covered under the Contract. w) **"Sexual Exploitation and Abuse" "(SEA)"** means the

			following: **"Sexual Exploitation"** is defined as means any actual or attempted abuse of position of vulnerability, differential power or trust, for sexual purposes, including, but not limited to, profiting monetarily, socially or politically from the sexual exploitation of another. **"Sexual Abuse"** is defined as means the actual or threatened physical intrusion of asexual nature, whether by force or under unequal or coercive conditions. x) **"Sexual Harassment" "(SH)"** is defined as means unwelcome sexual advances, requests for sexual favors, and other verbal or physical conduct of a sexual nature by the Contractor's Personnel with other Contractor's or Employer's Personnel. y) **"Sub-supplier"** means any person, whether natural or legal, named in the Contract as a Sub-contractor, or any person appointed as a Sub-contractor for a part of the Supplies and the legal successors in title to each of these persons. z) **"Supplier"** means the individual or firm supplying the Goods and Services under this Contract. Details of the supplier will be available in the Contract Finalization Information Section in the Contract Agreement. aa) **"Sustainable Public Procurement"** means a process whereby the Purchaser meets the need for goods, services and works in a way that achieves value for money on a whole lifecycle basis in terms of generating benefits not only to the Purchaser but also to society and the economy, whilst minimizing damage to the environment.
2.	Application and Interpretation	2.1	In interpreting these Conditions of Contract headings and marginal notes are used for convenience only and shall not affect their interpretations unless specifically stated; references to singular include the plural and vice versa;

			and masculine include the feminine. Words have their ordinary meaning under the language of the Contract unless specifically defined
		2.2	If any of the Contract Documents, correspondence or communications are prepared in any language other than the governing language under GCC 4.1 above, the English translation of such documents, correspondence or communications shall prevail in matters of interpretation.
		2.3	The documents forming the Contract shall be interpreted in the following order of priority: a) Form of Contract; b) Letter of Acceptance; c) Form of Tender and Priced Schedule; d) Special Conditions of Contract; e) General Conditions of Contract; f) Specifications; g) Performance Security; h) Power of Attorney; and i) Any other document listed in the **SCC** as forming part of the Contract Documents including Minutes of Negotiation, JV Agreement where applicable. [This should be shown as Appendices: From Appendix 1 - Appendix n^{th}
3.	Condition Precedent	3.1	The Contract shall come into effect after the Supplier fulfilling the conditions precedent stated in the **SCC**.
		3.2	If the Condition precedent stipulated on GCC 3.1 is not met by the date specified in the **SCC** this contract shall not come into effect;
		3.3	If the Purchaser is satisfied that each of the conditions precedent in this contract has been satisfied (except to the extent waived by him, but subject to such conditions as he

			shall impose in respect of such waiver) he shall promptly issue to the Supplier a certificate of Contract commencement, which shall confirm the start date.
4.	Governing Language	4.1	The Contract shall be written in the language specified in the **SCC**. Subject to this clause, the version of the Contract written in the specified language shall govern its interpretation. All correspondence and other documents pertaining to the Contract that are exchanged by the parties shall be written in the same language.
5.	Labor Law and other Applicable Laws	5.1	The Supplier shall comply with all the relevant labour laws applicable in the country, including laws relating to workers employment, working hours, welfare, immigration and anti- trafficking in persons.
		5.2	The Supplier shall require if any its Sub-Contractor and Personnel to obey all applicable laws, including those concerning safety at work.
		5.3	The Contract shall be interpreted in accordance with the laws of Tanzania.
6.	Country of Origin	6.1	All Goods and Services supplied under the Contract shall have their origin in the countries and territories eligible under projects financed by the specified institution, as further elaborated in the **SCC**.
		6.2	For purposes of this Clause, "**origin**" means the place where the Goods were mined, grown, or produced, or from which the Services are supplied. Goods are produced when, through manufacturing, processing, or substantial and major assembly of components, a commercially recognized new product results that is substantially different in basic characteristics or in purpose or utility from its components.
		6.3	The origin of Goods and Services may be distinct from the nationality of the Supplier.

7.	Standards	7.1	The Goods supplied under this Contract shall conform to the standards mentioned in the Technical Specifications, and, when no applicable standard is mentioned, to the authoritative standards appropriate to the Goods' country of origin. Such standards shall be the latest issued by the concerned institution.
8.	Use of Contract Documents and Information; Inspection and Audit by the Government of Tanzania	8.1	The Supplier shall not, without the Purchaser's prior written consent, disclose the Contract, or any provision thereof, or any specification, plan, drawing, pattern, sample, or information furnished by or on behalf of the Purchaser in connection therewith, to any person other than a person employed by the Supplier in the performance of the Contract. Disclosure to any such employed person shall be made in confidence and shall extend only so far as may be necessary for purposes of such performance.
		8.2	The Supplier shall not, without the Purchaser's prior written consent, make use of any document or information enumerated in GCC 7.1 [Standards] except for purposes of performing the Contract.
		8.3	Any document, other than the Contract itself, enumerated in GCC 7.1 [Standards] shall remain the property of the Purchaser and shall be returned (all copies) to the Purchaser on completion of the Supplier's performance under the Contract if so required by the Purchaser.
		8 .4	The Supplier shall permit the Government of Tanzania or / and donor agencies involved in financing the project to inspect the Supplier's accounts and records relating to the performance of the Supplier and to have them audited by auditors appointed by the Government of Tanzania or / and the appropriate donor agencies, if so required by the Government of Tanzania or / and the appropriate donor agencies.
9.	Certification	9.1	If required under the Applicable Law, Goods supplied

	of Goods in Accordance with Laws of the United Republic of Tanzania		under the Contract shall be registered for use in the United Republic of Tanzania as specified in the **SCC**. The Purchaser undertakes to cooperate with the Supplier to facilitate registration of the Goods for use in the United Republic of Tanzania.
		9.2	Unless otherwise specified in the **SCC**, the Contract shall become effective on the date ('the Effective Date") that the Supplier receives written notification from the relevant authority in the United Republic of Tanzania that the Goods have been registered for use in the United Republic of Tanzania.
		9.3	If thirty (30) days, or such longer period specified in the **SCC**, elapse from the date of Contract signing and the Contract has not become effective pursuant to GCC 9.2 above, then either party may, by not less than seven (7) days' written notice to the other party, declare this Contract null and void. In such event, the Supplier's performance security shall be promptly returned.
10.	Performance Securities	10.1	The Performance Securing declaration or Performance Security shall be provided to the Purchaser no later than the date specified in the Letter of Acceptance. In the case of Performance Security, it shall be issued in an amount and form and by a bank or surety acceptable to the Purchaser, and denominated in the types and proportions of the currencies in which the Contract Price is payable as specified in the **SCC**.
		10.2	The proceeds of the performance security shall be payable to the Purchaser as compensation for any loss resulting from the Supplier’s failure to complete its obligations under the Contract.
		10.3	The performance security will be discharged by the Purchaser and returned to the Supplier not later than thirty (30) days following the date of completion of the Supplier’s performance obligations under the Contract,

			including any warranty obligations, unless otherwise specified in **SCC**.
		10.4	Where circumstances necessitate the amendment of the contract after signature, and such amendment is effected, the Purchaser shall require the Supplier to provide additional Performance Security to cover any cumulative increase of more than ten percent for the Unconditional Bank Guarantee or 15% for Surety Bond of the initial Contract Price.
		10.5	In the case of Performance Securing Declaration, it shall remain in force until completion of the Supplier's performance obligations under the Contract, and in the event the Supplier failing to perform the obligations under the Contract, the Purchaser, following the termination of the contract, shall initiate the blacklisting process with the Public Procurement Regulatory Authority.
11.	Inspections and Tests	11.1	The Supplier shall at its own expense and at no cost to the Purchaser carry out all such tests and/or inspections of the Goods and Related Services as are specified in the **SCC**.
		11.2	The inspections and tests may be conducted on the premises of the Supplier or the manufacturer, at point of delivery, and/or at the Goods' final destination, or in another place in the Purchaser's Country as specified in the **SCC**. Subject to GCC11.3, if conducted on the premises of the Supplier or the manufacturer, all reasonable facilities and assistance, including access to production data, shall be furnished to the inspectors at no charge to the Purchaser
		11.3	The Purchaser or its designated representative shall be entitled to attend the tests and/or inspections referred to in GCC11.2, provided that the Purchaser bear all of its own costs and expenses incurred in connection with such attendance including, but not limited to, all traveling and board and lodging expenses.

		11.4	Whenever the Supplier is ready to carry out any such test and inspection, it shall give a reasonable advance notice, including the place and time, to the Purchaser. The Supplier shall obtain from any relevant third party or manufacturer any necessary permission or consent to enable the Purchaser or its designated representative to attend the test and/or inspection. (a) Said inspection and testing is for the Purchaser's account. In the event that inspection and testing is required prior to dispatch, the Goods shall not be shipped unless a satisfactory inspection and quality control report has been issued in respect of those Goods. (b) The Supplier may have an independent quality test conducted on a batch ready for shipment. The cost of such tests will be borne by the Supplier. (c) Upon receipt of the Goods at place of final destination, the Purchaser's representative shall inspect the Goods or part of the Goods to ensure that they conform to the condition of the Contract and advise the Purchaser that the Goods were received in apparent good order. The Purchaser will issue an Acceptance Certificate to the Supplier in respect of such Goods (or part of Goods). The Acceptance Certificate shall be issued within ten (10) days of receipt of the Goods or part of Goods at place of final destination
		11.5	Where the Supplier contests the validity of the rejection by the Purchaser or his representative, of any inspection as required by GCC11.4 above conducted before shipment or at ultimate destination, whether based on product or packing grounds, a sample drawn jointly by the Supplier and Purchaser or his or her representative and authenticated by both, will be forwarded for umpire analysis within four weeks of the time the Supplier contests to an independent agency mutually agreed by the Purchaser and Supplier. The umpire's finding, which will be promptly obtained, will be final and binding on both

			parties. The cost of umpire analysis will be borne by the losing party
		11.6	The Purchaser may require the Supplier to carry out any test and/or inspection not required by the Contract but deemed necessary to verify that the characteristics and performance of the Goods comply with the technical specifications' codes and standards under the Contract, provided that the Supplier's reasonable costs and expenses incurred in the carrying out of such test and/or inspection shall be added to the Contract Price. Further, if such test and/or inspection impedes the progress of manufacturing and/or the Supplier's performance of its other obligations under the Contract, due allowance will be made in respect of the Delivery Dates and Completion Dates and the other obligations so affected.
		11.7	The Supplier shall provide the Purchaser with a report of the results of any such test and/or inspection.
		11.8	The Purchaser may reject any Goods or any part thereof that fail to pass any test and/or inspection or do not conform to the specifications. The Supplier shall either rectify or replace such rejected Goods or parts thereof or make alterations necessary to meet the specifications at no cost to the Purchaser, and shall repeat the test and/or inspection, at no cost to the Purchaser, upon giving a notice pursuant to GCC 11.4.
		11.9	The Supplier agrees that neither the execution of a test and/or inspection of the Goods or any part thereof, nor the attendance by the Purchaser or its representative, nor the issue of any report pursuant to GCC 11.7, shall release the Supplier from any warranties or other obligations under the Contract.
12.	Packing	12.1	The Supplier shall provide such packing of the Goods as is required to prevent their damage or deterioration during transit to their final destination, as indicated in the

			Contract. The packing shall be sufficient to withstand, without limitation, rough handling during transit and exposure to extreme temperatures, salt, and precipitation during transit and open storage. Packing case size and weights shall take into consideration, where appropriate, the remoteness of the Goods' final destination and the absence of heavy handling facilities at all points in transit.
		12.2	The packing, marking, and documentation within and outside the packages shall comply strictly with such special requirements as shall be expressly provided for in the Contract, including additional requirements, if any, specified in the **SCC** or Technical Specifications, and in any subsequent instructions ordered by the Purchaser.
13.	Delivery and Documents	13.1	Delivery of the Goods shall be made by the Supplier in accordance with the terms specified in the Schedule of Requirements. The details of shipping and/or other documents to be furnished by the Supplier are specified in the **SCC**.
		13.2	For purposes of the Contract, "EXW," "FOB," 'FCA,' 'CIF," "CIP," and other trade terms used to describe the obligations of the parties shall have the meanings assigned to them by the current edition of Incoterms published by the International Chamber of Commerce, Paris.
		13.3	Documents to be submitted by the Supplier are specified in the **SCC**.
14.	Insurance	14.1	The Goods supplied under the Contract shall be fully insured in a freely convertible currency against loss or damage incidental to manufacture or acquisition, transportation, storage, and delivery in the manner specified in the **SCC**.
		14.2	Where delivery of the Goods is required by the Purchaser on a CIF or CIP basis, the Supplier shall arrange and pay for cargo insurance, naming the Purchaser as beneficiary.

			Where delivery is on an FOB or FCA basis, insurance shall be the responsibility of the Purchaser.
15.	Transportation	15.1	Where the Supplier is required under Contract to deliver the Goods FOB, transport of the Goods, up to and including the point of putting the Goods on board the vessel at the specified port of loading, shall be arranged and paid for by the Supplier, and the cost thereof shall be included in the Contract Price. Where the Supplier is required under the Contract to deliver the Goods FCA, transport of the Goods and delivery into the custody of the carrier at the place named by the Purchaser or other agreed point shall be arranged and paid for by the Supplier, and the cost thereof shall be included in the Contract Price.
		15.2	Where the Supplier is required under Contract to deliver the Goods CIF or CIP, transport of the Goods to the port of destination or such other named place of destination in the United Republic of Tanzania, as shall be specified in the Contract, shall be arranged and paid for by the Supplier, and the cost thereof shall be included in the Contract Price.
		15.3	Where the Supplier is required under the Contact to transport the Goods to a specified place of destination within the United Republic of Tanzania, defined as the Site, transport to such place of destination in the United Republic of Tanzania, including insurance and storage, as shall be specified in the Contract, shall be arranged by the Supplier, and related costs shall be included in the Contract Price.
		15.4	Where the Supplier is required under Contract to deliver the Goods CIF or CIP, no restriction shall be placed on the choice of carrier. Where the Supplier is required under Contract (a) to deliver the Goods FOB or FCA, and (b) to arrange on behalf and at the expense of the Purchaser for international transportation on specified carriers or on

			national flag carriers of the United Republic of Tanzania, the Supplier may arrange for such transportation on alternative carriers if the specified or national flag carriers are not available to transport the Goods within the period(s) specified in the Contract.
16.	Incidental Services	16.1	The Supplier shall provide such incidental services, if any, as are specified in the **SCC**.
		16.2	Prices charged by the Supplier for incidental services, if not included in the Contract Price for the Goods, shall be agreed upon in advance by the parties and shall not exceed the prevailing rates charged to other parties by the Supplier for similar services.
17.	Spare Parts	17.1	As specified in **SCC**, the Supplier may be required to provide any or all of the following materials, notifications, and information pertaining to spare parts manufactured or distributed by the Supplier: (a) Such spare parts as the Purchaser may elect to purchase from the Supplier, provided that this election shall not relieve the Supplier of any warranty obligations under the Contract; and (b) In the event of termination of production of the spare parts: (i) advance notification to the Purchaser of the pending termination, in sufficient time to permit the Purchaser to procure needed requirements; and (ii) following such termination, furnishing at no cost to the Purchaser, the blueprints, drawings, and specifications of the spare parts, if requested.
18.	Warranty	18.1	All goods must be of fresh manufacture and must bear the dates of manufacture and expiry. The Supplier further warrants that all Goods supplied under the Contract will have remaining a minimum of five-sixths (5/6) of the specified shelf life upon delivery at

			port/airport of entry for goods with a shelf life of more than two years and three-fourths (3/4) for goods with a shelf life of two years or less, unless otherwise specified in the **SCC**; have "averages" within the ranges set forth in the Technical Specifications, where applicable; are not subject to recall by the applicable regulatory authority due to unacceptable 'quality or an adverse drug reaction; and in every other respect will fully comply in all respects with the Technical Specifications and with the conditions laid down in the Contract.
		18.2	The Purchaser shall have the right to make claims under the above warranty for the period specified in the **SCC** after the Goods have been delivered to the final destination indicated in the Contract. Upon receipt of a written notice from the Purchaser, the Supplier shall, with all reasonable speed, replace the defective Goods without cost to the Purchaser. The Supplier will be entitled to remove, at his own risk and cost, the defective Goods once the replacement Goods have been delivered.
		18.3	In the event of a dispute by the Supplier, a counter-analysis will be carried out on the manufacturer's retained samples by an independent neutral laboratory agreed by both the Purchaser and the Supplier. If the counter-analysis confirms the defect, the cost of such analysis will be borne by the Supplier as well as the replacement and disposal of the defective goods. In the event of the independent analysis confirming the quality of the product, the Purchaser will meet all costs for such analysis.
		18.4	If, after being notified that the defect has been confirmed pursuant to GCC 18.2 above, the Supplier fails to replace the defective Goods within the period specified in the **SCC**, the Purchaser may proceed to take such remedial action as may be necessary, including removal and disposal, at the Supplier's risk and expense and without

			prejudice to any other rights that the Purchaser may have against the Supplier under the Contract. The Purchaser will also be entitled to claim for storage in respect of the defective Goods for the period following notification and deduct the sum from payments due to the Supplier under this Contract.
		18.5	In the event any of the Goods are recalled, the Supplier shall notify the Purchaser within fourteen (14) days, providing full details of the reason for the recall and promptly replace, at its own cost, the items covered by the recall with Goods that fully meet the requirements of the Technical Specification and arrange for collection or destruction of any defective Goods. If the Supplier fails to fulfil its recall obligation promptly, the Purchaser will, at the Supplier's expense, carry out the recall.
19.	Patent and Copy Rights	19.1	The Supplier shall, subject to the Purchaser's compliance with GCC18.2, indemnify and hold harmless the Purchaser and its employees and officers from and against any and all suits, actions or administrative proceedings, claims, demands, losses, damages, costs, and expenses of any nature, including attorney's fees and expenses, which the Purchaser may suffer as a result of any infringement or alleged infringement of any patent, utility model, registered design, trademark, copyright, or other intellectual property right registered or otherwise existing at the date of the Contract by reason of: a) the installation of the Goods by the Supplier or the use of the Goods in the country where the Site is located; and b) the sale in any country of the products produced by the Goods. Such indemnity shall not cover any use of the Goods or any part thereof other than for the purpose indicated by or to be reasonably inferred from the Contract, neither any infringement resulting from the use of the Goods or any part thereof, or any products produced thereby in

			association or combination with any other equipment, plant, or materials not supplied by the Supplier, pursuant to the Contract.
		19.2	If any proceedings are brought or any claim is made against the Purchaser arising out of the matters referred to in GCC18.1, the Purchaser shall promptly give the Supplier a notice thereof, and the Supplier may at its own expense and in the Purchaser's name conduct such proceedings or claim and any negotiations for the settlement of any such proceedings or claim.
		19.3	If the Supplier fails to notify the Purchaser within twenty-eight (28) days after receipt of such notice that it intends to conduct any such proceedings or claim, then the Purchaser shall be free to conduct the same on its own behalf.
		19.4	The Purchaser shall, at the Supplier's request, afford all available assistance to the Supplier in conducting such proceedings or claim, and shall be reimbursed by the Supplier for all reasonable expenses incurred in so doing.
		19.5	The Purchaser shall indemnify and hold harmless the Supplier and its employees, officers, and Subcontractors from and against any and all suits, actions or administrative proceedings, claims, demands, losses, damages, costs, and expenses of any nature, including attorney's fees and expenses, which the Supplier may suffer as a result of any infringement or alleged infringement of any patent, utility model, registered design, trademark, copyright, or other intellectual property right registered or otherwise existing at the date of the Contract arising out of or in connection with any design, data, drawing, specification, or other documents or materials provided or designed by or on behalf of the Purchaser.
20.	Limitation of	20.1	Except in cases of criminal negligence or wilful

	Liability		misconduct, and in the case of infringement pursuant to Clause 18, a) the Supplier shall not be liable to the Purchaser, whether in contract, tort, or otherwise, for any indirect or consequential loss or damage, loss of use, loss of production, or loss of profits or interest costs, provided that this exclusion shall not apply to any obligation of the Supplier to pay liquidated damages to the Purchaser; and b) the aggregate liability of the Supplier to the Purchaser, whether under the Contract, in tort or otherwise, shall not exceed the total Contract Price, provided that this limitation shall not apply to the cost of repairing or replacing defective equipment, or to any obligation of the supplier to indemnify the Purchaser with respect to patent infringement.
21.	Change of Laws and Regulations	21.1	Unless otherwise specified in the Contract, if after the date of 28 days prior to the deadline date of Tender submission, any law, regulation, ordinance, order or bylaw having the force of law is enacted, promulgated, abrogated, or changed in the place of the Purchaser's Country where the Site is located (which shall be deemed to include any change in interpretation or application by the competent authorities) that subsequently affects the Delivery Date and/or the Contract Price, then such Delivery Date and/or Contract Price shall be correspondingly increased or decreased, to the extent that the Supplier has thereby been affected in the performance of any of its obligations under the Contract. Notwithstanding the foregoing, such additional or reduced cost shall not be separately paid or credited if the same has already been accounted for in the price adjustment provisions where applicable, in accordance with GCC 23 [Prices].
22.	Payment	22.1	The method and conditions of payment to be made to the Supplier under this Contract shall be specified in the **SCC**.

		22.2	The Supplier's request(s) for payment shall be made to the Purchaser in writing or in electronic forms that provide record of the content of communication, accompanied by an invoice describing, as appropriate, the Goods delivered and Services performed, and by documents submitted pursuant to GCC 13 [Delivery and Documents], and upon fulfilment of other obligations stipulated in the Contract.
		22.3	Payments shall be made promptly by the Purchaser, in the mode of payment specified in the **SCC**, but in no case later than sixty (60) days after submission of an invoice or claim by the Supplier. In the event that the Purchaser fails to make any payment by its respective due date or within the period set forth in the Contract, the Purchaser shall pay to the Supplier interest on the amount of such delayed payment at the rate(s) shown in SCC to the Agreement for the period of delay until payment has been made in full, whether before or after judgment or arbitrage award. There shall be a cumulative cap on the interest charges of up to 10% of the delayed payment at which point the Supplier may consider termination of the Contract.
		22.4	The currency or currencies in which payment is made to the Supplier under this Contract shall be specified in the **SCC** subject to the following general principle: Payment will be made in the currency or currencies in which the payment has been requested in the Supplier's tender.
23.	Prices	23.1	The contract price shall be as specified in the Contract Agreement Subject to any additions and adjustments thereto or deductions there from, as may be made pursuant to the Contract.
		23.2	Prices charged by the Supplier for Goods delivered and Services performed under the Contract shall not vary from the prices quoted by the Supplier in its tender, with the exception of any price adjustments authorized in the **SCC**

			or in the Purchaser's request for tender validity extension, as the case may be.
		23.3	Prices payable to the Supplier, if subject to adjustment during performance of the Contract to reflect changes in the cost of labour and material components shall be done in accordance with GCC 23.4.
		23.4	If in accordance with GCC 23.2 and 23.3, prices shall be adjustable, the following method shall be used to calculate the price adjustment: Prices payable to the Supplier, as stated in the Contract, shall be subject to adjustment during performance of the Contract to reflect changes in the cost of labour and material components in accordance with the formula: P1 = P0 [a + bL1/L0 +cM1/M0] - P0 where a + b + c = 1 in which: P1 = adjustment amount payable to the Supplier. P0 = Contract Price(base price). a = fixed element representing profits and overheads included in the Contract Price and generally in the range of five (5) to fifteen (15) percent. b = estimated percentage of labour component in the Contract Price. c = estimated percentage of material component in the Contract Price. L0, L1 = labour indices applicable to the appropriate industry in the country of origin on the base date and date for adjustment, respectively. M0, M1 = material indices for the major raw material on the base date and date for adjustment, respectively, in the country of origin. The Bidder shall indicate the source of the indices and the

			base date indices in its bid. The coefficients a, b, and c as specified by the Purchaser are as follows: a =[insert value of coefficient] b= [insert value of coefficient] c= [insert value of coefficient] Base date = thirty (30) days prior to the deadline for submission of the bids. Date of adjustment =mid-point of the period of manufacture). The above price adjustment formula shall be invoked by either party subject to the following further conditions: (a) No price adjustment shall be allowed beyond the original delivery dates. As a rule, no price adjustment shall be allowed for periods of delay for which the Supplier is entirely responsible. The Purchaser will, however, be entitled to any decrease in the prices of the Goods and Services subject to adjustment. (b) If the currency in which the Contract Price P0 is expressed is different from the currency of origin of the labour and material indices, a correction factor will be applied to avoid incorrect adjustments of the Contract Price. The correction factor shall be: Z0/Z1, where: Z0 =the number of units of currency of the origin of the indices which equal to one unit of the currency of the Contract Price on the Base date, and Z1= the number of units of currency of origin of the indices which equal to one unit of the currency of the Contract Price on the Date for Adjustment. (c) No price adjustment shall be payable on the portion of the Contract Price paid to the Supplier as advance payment.
24.	Change	24.1	The estimated quantities required are shown in the

	Orders, Quantities and Change Orders		schedule of requirements. Although every endeavor has been made to prepare an accurate estimate of requirements during the period of the Contract, the Purchaser does not bind himself to take the quantities stated, but reserves the right to order greater or lesser of quantities according to the requirements during that time.
		24.2	The Purchaser may at any time, by a written order given to the Supplier make changes within the general scope of the Contract and any one or more of the following: a) Drawings, designs, or specifications, where Goods to be furnished under the Contract are to be specifically manufactured for the Purchaser; b) The method of shipment or packing; c) The place of delivery; and / or d) The Services to be provided by the Supplier.
		24.3	If any such change causes an increase or decrease in the cost of, or the time required for, the Supplier's performance of any provisions under the Contract, an equitable adjustment shall be made in the Contract, an equitable adjustment shall be made in the Contract Price or delivery schedule, or both, and the Contract shall accordingly be amended. Any claims by the Supplier for adjustment under this clause must be asserted within thirty (30) days from the date of the Supplier's receipt of the Purchaser's change order.
		24.4	Prices to be charged by the supplier for any related services that might be needed but which were not included in the Contract shall be agreed upon in advance by the Parties and shall not exceed the prevailing rates charged to other parties by the Supplier for similar services.
25.	Extensions of Time	25.1	If at any time during performance of the Contract, the Supplier or its subcontractors should encounter conditions impeding timely delivery of the Goods or completion of

			Related Services pursuant to GCC13 [Delivery and Documents], the Supplier shall promptly notify the Purchaser in writing of the delay, its likely duration, and its cause. As soon as practicable after receipt of the Supplier's notice, the Purchaser shall evaluate the situation and may at its discretion extend the Supplier's time for performance, in which case the extension shall be ratified by the parties by amendment of the Contract.
		25.2	Except in case of Force Majeure, as provided under GCC31, a delay by the Supplier in the performance of its Delivery and Completion obligations shall render the Supplier liable to the imposition of liquidated damages pursuant to GCC 29 [Liquidated Damages], unless an extension of time is agreed upon, pursuant to GCC 25.1.
26.	Assignment	26.1	The Supplier shall not assign, in whole or in part, its obligations to perform under this Contract, except with the Purchaser's prior written consent.
27.	Subcontracting	27.1	The Supplier shall consult the Purchaser in the event of subcontracting under this contract if not already specified in the Tender. Subcontracting shall not alter the Supplier's obligations.
28.	Delays in the Supplier's Performance	28.1	Delivery of the Goods and performance of Services shall be made by the Supplier in accordance with the time schedule prescribed by the Purchaser in the Schedule of Requirements.
		28.2	If at any time during performance of the Contract, the Supplier or its subcontractor(s) should encounter conditions impeding timely delivery of the Goods and performance of Services, the Supplier shall promptly notify the Purchaser in writing or in electronic forms that provide record of the content of communication of the fact of the delay, its likely duration, and its cause(s). As soon as practicable after receipt of the Supplier's notice, the Purchaser shall evaluate the situation and may at its

			discretion extend the Supplier's time for performance, with or without liquidated damages, in which case the extension shall be ratified by the parties by amendment of Contract.
		28.3	Except as provided under GCC 31 [Force Majeure], a delay by the Supplier in the performance of its delivery obligations shall render the Supplier liable to the imposition of liquidated damages pursuant to GCC 29 [Liquidated Damages}, unless an extension of time is agreed upon pursuant to GCC 25.2 without the application of liquidated damages.
29.	Liquidated Damages	29.1	Subject to GCC 31 [Force Majeure], if the Supplier fails to deliver any or all of the Goods or to perform the Services within the period(s) specified in the Contract, the Purchaser shall, without prejudice to its other remedies under the Contract, deduct from the Contract Price, as liquidated damages, a sum equivalent to the percentage specified in the **SCC** of the delivered price of the delayed Goods or unperformed Services for each week or part thereof of delay until actual delivery or performance, up to a maximum deduction of the percentage specified in the **SCC**. Once the maximum is reached, the Purchaser may consider termination of 30 [Termination for Default].
30.	Termination for Default	30.1	The Purchaser, without prejudice to any other remedy for breach of Contract, by written notice of default sent to the Supplier, may terminate this Contract in whole or in part: a) if the Supplier fails to deliver any or all of the Goods within the period(s) specified in the Contract, or within any extension thereof granted by the Purchaser pursuant to GCC 28 [Delays in the Supplier's Performance]; or b) if the Goods do not meet the Technical Specifications stated in the Contract; or c) if the Supplier fails to provide any registration or other certificates in respect of the Goods within the time

			specified in the Special Conditions; or d) if the Supplier, in the judgment of the Purchaser, has engaged in corrupt or fraudulent practices in competing for or in executing the Contract. For the purpose of this clause: "**corrupt practice**" means the offering, giving receiving or soliciting of anything of value to influence the action of a public officer in the procurement process or contract execution. "**coercive practice**" means impairing or harming, or threatening to impair or harm directly or indirectly, any party or the property of the party for the purpose of influencing improperly the action or that party in connection with public procurement or in furtherance of corrupt practice or fraudulent practice.
		30.2	In the event the Purchaser terminates the Contract in whole or in part, pursuant to GCC 30.1, the Purchaser may procure, upon such terms and in such manner as it deems appropriate, Goods or Services similar to those undelivered, and the Supplier shall be liable to the Purchaser for any excess costs for such similar Goods or Services. However, the Supplier shall continue performance of the Contract to the extent not terminated.
31.	Force Majeure	31.1	Notwithstanding the provisions of GCC 28 [Delays in Supplier Performance], GCC 29 [Liquidated Damages] and GCC 30 [Termination for Default], the Supplier shall not be liable for forfeiture of its performance security, liquidated damages, or termination for default if and to the extent that its delay in performance or other failure to perform its obligations under the Contract is the result of an event of Force Majeure.
		31.2	If a Force Majeure situation arises, the Supplier shall promptly notify the Purchaser in writing or in electronic forms that provide record of the content of communication

			of such condition and the cause thereof. Unless otherwise directed by the Purchaser in writing, the Supplier shall continue to perform its obligations under the Contract as far as is reasonably practical and shall seek all reasonable alternative means for performance not prevented by the Force Majeure event.
32.	Termination for Insolvency	32.1	The Purchaser may at any time terminate the Contract by giving written notice to the Supplier if the Supplier becomes bankrupt or otherwise insolvent. In this event, termination will be without compensation to the Supplier, provided that such termination will not prejudice or affect any right of action or remedy that has accrued or will accrue thereafter to the Purchaser.
33.	Termination for Convenience	33.1	The Purchaser, by written notice sent to the Supplier, may terminate the Contract, in whole or in part, at any time for its convenience. The notice of termination shall specify that termination is for the Purchaser's convenience, the extent to which performance of the Supplier under the Contract is terminated, and the date upon which such termination becomes effective.
		33.2	The Goods that are complete and ready for shipment within thirty (30) days after the Supplier's receipt of notice of termination shall be accepted by the Purchaser at the Contract terms and prices. For the remaining Goods, the Purchaser may elect: a) to have any portion completed and delivered at the Contract terms and prices; and/or b) to cancel the remainder and pay to the Supplier an agreed amount for partially completed Goods and Services and for materials and parts previously procured by the Supplier.
34.	Settlement of Disputes	34.1	In the event of any dispute arising out of this contract, either party shall issue a notice of dispute to settle the dispute amicably. The parties hereto shall, within twenty

			eight (28) days from the notice date, use their best efforts to settle the dispute amicably through mutual consultations and negotiation.
		34.2	If, after fourteen (14) days, the parties have failed to resolve their dispute or difference by such mutual consultation, then either the Supplier or the Purchaser may give notice to the other party of its intention to commence arbitration, as hereinafter provided, as to the matter in dispute, and no arbitration in respect of this matter may be commenced unless such notice is given. Any dispute or difference in respect of which a notice of intention to commence arbitration has been given in accordance with this Clause shall be finally settled by arbitration. Arbitration may be commenced prior to or after handover of the Assets under the Contract.
		34.3	The arbitration shall be conducted in accordance with the arbitration procedure published by the Institution named and, in the place, shown in the **SCC**.
		34.4	Notwithstanding any reference to arbitration herein; a) the parties shall continue to perform their respective obligations under the Contract unless they otherwise agree; and b) The Purchaser shall pay the Supplier any monies due the Supplier.
35.	Notices	35.1	Any notice given by one party to the other pursuant to this Contract shall be sent to the other party in writing or in electronic forms that provide record of the content of communication and confirmed in writing or in electronic forms that provide record of the content of communication to the other party's address as indicated in the **SCC.** Details of the supplier' address will be available in the Contract Finalization Information Section in the Contract Agreement.

		35.2	A notice shall be effective when delivered or on notice's effective date, whichever is later.
36.	Taxes and Duties	36.1	A Supplier supplying Goods from abroad shall be entirely responsible for all taxes, stamp, duties, license fees, and other such levies imposed outside the United Republic of Tanzania.
		36.2	A Supplier supplying Goods offered locally shall be entirely responsible for all taxes, duties, license fees, etc., incurred until delivery of the contracted Goods the Purchaser.
		36.3	If any tax exemptions, reductions, allowances or privileges may be available to the Supplier in the United Republic of Tanzania the Purchaser shall use its best efforts to enable the Supplier to benefit from any such tax savings to the maximum allowable extent.
37.	Suspension of Financing	37.1	In the event that the source of financing is suspended to the Purchaser, from which part of the payments to the Contractor are being made: a) The Purchaser is obligated to notify the Contractor of such suspension within 7 days of having received the financing agency's suspension notice. b) If the Contractor has not received sums due it within the 28 days for payment provided for in GCC 22.3 [Payment], the Contractor may immediately issue a 14 days termination notice.
38.	Delayed Payments	38.	Unless otherwise specified in the SCC, the event of a late payment by the Purchaser, the Supplier shall be entitled to an interest on the delayed payment, which will be included in the next payment. Interest shall be calculated from the due date of the payment until the actual date of payment, at the prevailing rate issued by the Bank of Tanzania (BOT) as of the Contract signing date, for each currency involved (domestic or foreign).The interest rate shall be

			the prevailing overall time deposit rate issued by BOT for each currency of payment as specified in the SCC. If negotiated, this rate shall not exceed 1% above the BOT prevailing rate. However, interest on delayed payments shall not exceed 10% of the unpaid amount and at this point the Supplier may consider terminating the Contract.

SECTION IX: SPECIAL CONDITIONS OF CONTRACT (SCC)

Special Conditions of Contract (SCC)

The following Special Conditions of Contract (SCC) shall supplement the General Condition Contract (SCC). Whenever there is a conflict, the provisions herein shall prevail over those in the GCC. The corresponding clause number of the GCC is indicated in parentheses.

SCC Number	Information/Data Required	GCC Number	Amendments of, and Supplements to, Clauses in the GCC
1.	Appointing Authority for the Arbitrator	1.1(a)	**Tanzania Institute of Arbitrators (TIArb)**
2.	End User	1.1 (k)	**DISTRICT MEDICAL OFFICER** **MBEYA REGION**
3.	Final Destination	1.1 (l)	**DISTRICT MEDICAL OFFICE IN ALL COUNCILS**
4.	Name of Purchaser	1.1 (r)	**MBEYA REGIONAL SECRETARIAT** **P.O BOX 754, MBEYA**
5.	Other Documents Forming the Contract	2.3 (i)	Other documents forming part of the contract are BRELA,TIN CERTIFICATE,TMDA AND PHARMANCY COUNCIL REGISTRATION,BUSINESS PERMIT,BUSINESS LICENSE, SUPERINTENDENT AGREEMENT,VALID PROFESSIONAL REGISTRATION CERTIFICATE AND LIECENSES.
6.	Conditions Precedent	3.1	APPLICABLE
7.	Date for fulfilling Conditions Precedent	3.2	Date for meeting Condition precedent shall be 14 days from the date of signing the contract.

8.	Governing Language	4.1	**ENGLISH**
9.	Country of Origin	6.1	Not Applicable
10.	Certification of Goods	9.1	**NONE**
11.	Effectiveness of Contract	9.2	**1095** days after signing the contract.
12.	Voidance of contract due to ineffectiveness	9.3	**14** days after contract signing.
13.	Performance Security/Performance Securing Declaration	10.1	Performance Securing Declaration shall be applicable.
13.	Discharge of Performance Security	10.3	Not Applicable
14.	Inspection and Tests	11.1 & 11.2	**LOT NO. 78/2025/2026/G/72** **Procurement of medicines from prime vendor for Councils health facilities** **GFS Code: 22004102 - Drugs and Medicines** • **CEPHALEXIN 250MG/5ML ORAL SUSPESNSION, 100MLS** **This item does not require pre-shipment inspection** • **LOPERAMIDE 2MG TABLETS (LOPA)** **This item does not require pre-shipment inspection** • **HEPARIN 1000IU/ML, 5ML INJ**

			This item does not require pre-shipment inspection • **DEXAMETHASONE 0.1% 5ML EYE DROPS (XSONE)** **This item does not require pre-shipment inspection** • **IBUPROFEN 200MG TABLETS** **This item does not require pre-shipment inspection** • **CLARITHROMYCIN 250MG TABLET** **This item does not require pre-shipment inspection** • **CHLORPHENIRAMINE 4MG TABLETS** **This item does not require pre-shipment inspection** • **CLONIDINE 300MCG TABLET** **This item does not require pre-shipment inspection** • **ETAMSYLATE 500MG TABLET.** **This item does not require pre-shipment inspection** • **CEFUROXIME 250MG TABS (THEOROXIME)** **This item does not require pre-shipment inspection** • **PIROXICAM 20MG CAPSULE** **This item does not require pre-shipment inspection**

			• **GLIMEPIRIDE 2+ METFORMIN 500MG TABLETS (ILET B2)** **This item does not require pre-shipment inspection** • **METFORMIN 500MG TABLETS** **This item does not require pre-shipment inspection** • **ATORVASTATIN 40MG TABLETS** **This item does not require pre-shipment inspection** • **CETIRIZINE TABLET 10MG (ALLERIEVE)** **This item does not require pre-shipment inspection** • **DIGOXIN 0.25MG TABLETS** **This item does not require pre-shipment inspection** • **ACETYLSALICYLIC ACID (ASPIRIN) 300MG TABLET** **This item does not require pre-shipment inspection** • **SALBUTAMOL 100MCG INHALER** **This item does not require pre-shipment inspection** • **BORIC ACID EAR DROPS** **This item does not require pre-shipment inspection** • **TAMSULOSIN 0.4MG TABLETS** **This item does not require pre-shipment inspection**

			• **CIPROFLOXACIN 500MG+ TINIDAZOLE 600MG TABLETS** **This item does not require pre-shipment inspection** • **PARACETAMOL 1000MG INJECTION (LENIOL)** **This item does not require pre-shipment inspection** • **CHLORPROMAZINE 100MG TABLETS** **This item does not require pre-shipment inspection** • **MAGNESIUM SULPHATE 250MG/ML (50%) 10MLS SOLUTION FOR INJECTION** **This item does not require pre-shipment inspection** • **METFORMIN 500MG + GLIBLENCLAMIDE 5MG TABLET** **This item does not require pre-shipment inspection** • **TELMISARTAN 80MG + HYDROCHLOROTHIAZIDE 12.5MG TABLETS (SAFETELMI-H)** **This item does not require pre-shipment inspection** • **ACTINOMYCINE D 0.5 MG INJECTION** **This item does not require pre-shipment inspection** • **ASCORBIC ACID 100MG TABLETS**

			This item does not require pre-shipment inspection • **NIFEDIPINE 20MG TABLETS** **This item does not require pre-shipment inspection** • **DICLOFENAC 50 MG + PARACETAMOL 500 MG TABLET** **This item does not require pre-shipment inspection** • **PROMETHAZINE HCL 5MG/5ML ORAL LIQUID, 60ML** **This item does not require pre-shipment inspection** • **BETAMETHASONE + SALICYLIC OINTMENT, 15GM** **This item does not require pre-shipment inspection** • **TIZANIDINE 4MG TABLETS** **This item does not require pre-shipment inspection** • **SODIUM CROMOGLYCATE 2% EYE DROPS, 10MLS** **This item does not require pre-shipment inspection** • **ARTESUNATE 60MG POWDER FOR INJECTION** **This item does not require pre-shipment inspection** • **HYDROCORTISONE 100MG, POWDER FOR INJECTION**

			This item does not require pre-shipment inspection • **CLOTRIMAZOLE PESSARY WITH APPLICATOR 100 MG** **This item does not require pre-shipment inspection** • **SILVER NITRATE STICK** **This item does not require pre-shipment inspection** • **BETAMETHASONE 0.025%+CLOTRIMAZOLE 1%+GENTAMICIN0.5% (SKDERM) 30GM** **This item does not require pre-shipment inspection** • **HYDROCORTISONE ACETATE EYE DROPS 0.5%** **This item does not require pre-shipment inspection** • **ARTEMETHER/LUMEFANTRINE 20/120 MG TABLETS** **This item does not require pre-shipment inspection** • **FLUCLOXACILLIN 125MG +AMOXICILLIN 125M, 250MG/5ML , 100ML** **This item does not require pre-shipment inspection** • **GLUCOSAMINE SULPHATE 500MG+ CHONDROITIN SULPHATE 400MG TABLETS**

			This item does not require pre-shipment inspection • **AZITHROMYCIN 500MG TABLET** **This item does not require pre-shipment inspection** • **BRIMONIDINE EYE DROP 0.2%, 10 MLS (BRIZAGAN)** **This item does not require pre-shipment inspection** • **CANDESARTAN TABLETS 16MG (CANDEZ 16)** **This item does not require pre-shipment inspection** • **MUPIROCIN 2% CREAM, 15GM** **This item does not require pre-shipment inspection** • **ANTI - RABIES IMMUNOGLOBULIN 1000IU/5MLS INJECTION** **This item does not require pre-shipment inspection** • **TRANEXAMIC ACID 500MG TABLET** **This item does not require pre-shipment inspection** • **GENTAMYCIN SULPHATE EYE DROPS, 0.3%, 10MLS** **This item does not require pre-shipment inspection** • **CLOBETASOL PROPIONATE 0.05% OINTMENT, 20GM**

			This item does not require pre-shipment inspection • **INSUIN MIXTARD (NOVOMIX) 3ML** **This item does not require pre-shipment inspection** • **OXYBUTYNIN 5MG CAPSULE** **This item does not require pre-shipment inspection** • **ESOMEPRAZOLE 40MG TABLETS** **This item does not require pre-shipment inspection** • **FLUCLOXACILLIN 250MG+ AMOXYCILLIN 250MG CAPSULES** **This item does not require pre-shipment inspection** • **AMPICILLIN 250MG+ CLOXACILLIN 250MG CAPSULES** **This item does not require pre-shipment inspection** • **AZITHROMYCIN DEHYDRATE 250MG TABLETS** **This item does not require pre-shipment inspection** • **INSULIN HUMAN SOLUBLE VIAL 100IU/ML, 10 ML INJECTION** **This item does not require pre-shipment inspection** • **DEXAMETHASONE 4MG TABLETS** **This item does not require pre-shipment inspection**

			• **CO-TRIMOXAZOLE 480MG TABLETS** **This item does not require pre-shipment inspection** • **AMOXICILLIN 125MG DISPERSIBLE TABLETS** **This item does not require pre-shipment inspection** • **FUSIDIC ACID CREAM 15G** **This item does not require pre-shipment inspection** • **FLUCONAZOLE 2MG/ ML,100MLS INJECTION** **This item does not require pre-shipment inspection** • **LINDANE (1%)+BENZOCAINE (2%) CREAM 25G** **This item does not require pre-shipment inspection** • **CEFTRIAXONE 1000 MG +SULBACTAM 500 MG POWDER FOR INJECTION (NEOMICEF)** **This item does not require pre-shipment inspection** • **VITAMIN A - RETINOL CAPSULES 200,000 IU CAPSULES** **This item does not require pre-shipment inspection** • **PHENOBARBITAL 30MG TABLETS** **This item does not require pre-shipment inspection**

			• **ALLOPURINOL 300MG TABLET** **This item does not require pre-shipment inspection** • **DEXAMETHASONE 4 MG/ML INJECTION** **This item does not require pre-shipment inspection** • **CAPTOPRIL 25MG TABLETS** **This item does not require pre-shipment inspection** • **VITAMIN A (RETINOL) 100,000 I.U CAPSULES** **This item does not require pre-shipment inspection** • **BENZOYL PEROXIDE 5% GEL, 20GM** **This item does not require pre-shipment inspection** • **OXYTETRACYCLINE 1% EYE OINTMENT** **This item does not require pre-shipment inspection** • **CLOMIPHENE 50MG TABLET** **This item does not require pre-shipment inspection** • **MULTIVITAMIN COMBINATION TABLET** **This item does not require pre-shipment inspection** • **SULPHADOXINE 500MG**

			+PYRIMETHAMINE 25MG TABLETS **This item does not require pre-shipment inspection** • **TRAMADOL 50MG CAPSULES** **This item does not require pre-shipment inspection** • **CRESOL SAPONATED (LYSOL) 50% LIQUID** **This item does not require pre-shipment inspection** • **HYDRALAZINE 20MG/ML 1ML INJECTION** **This item does not require pre-shipment inspection** • **ACYCLOVIR EYE OINTMENT** **This item does not require pre-shipment inspection** • **FEBUXOSTAT 40MG TABLETS** **This item does not require pre-shipment inspection** • **TERBINAFINE 15G CREAM 1%** **This item does not require pre-shipment inspection** • **LATANOPROST 0.005% EYE SOLUTION** **This item does not require pre-shipment inspection** • **ERYTHROMYCIN 250MG TABLETS** **This item does not require pre-shipment inspection**

			• **METRONIDAZOLE 200MG TABLETS** **This item does not require pre-shipment inspection** • **MEBENDAZOLE SUSPENSION 100MG/5ML** **This item does not require pre-shipment inspection** • **FLUDROCORTISONE 0.1MG TABLETS** **This item does not require pre-shipment inspection** • **SODIUM LACTATE COMPOUND,500MLS ,SOLUTION FOR INTRAVENOUS INFUSSION** **This item does not require pre-shipment inspection** • **MUPIROCIN 2% OINTMENT 15GM** **This item does not require pre-shipment inspection** • **CHLORHEXIDINE 1.5% + CETRIMIDE 15% LIQUID** **This item does not require pre-shipment inspection** • **CLONAZEPAM 2MG TABLET** **This item does not require pre-shipment inspection** • **CALAMINE LOTION, 100ML** **This item does not require pre-shipment inspection**

			• **KETAMINE 10ML 50MG/ML INJECTION.** **This item does not require pre-shipment inspection** • **LINEZOLID 400MG TABLET** **This item does not require pre-shipment inspection** • **MICONAZOLE 40GM VAGINAL CREAM (GYNAZOL)** **This item does not require pre-shipment inspection** • **BETAMETHASONE (VALERATE) OINTMENT OR CREAM 0.025%** **This item does not require pre-shipment inspection** • **BETAMETHASONE + GENTAMYCIN EYE DROPS** **This item does not require pre-shipment inspection** • **ISOFLURANE INHALATION 250ML BOTTLE** **This item does not require pre-shipment inspection** • **FERROUS FUMARATE/ANTI ANAEMIA SYRUPS 20MG/ML, 200 MLS (HEMOVIT)** **This item does not require pre-shipment inspection** • **FLUPHENAZINE DECANOATE 25 MG/ML INJECTION** **This item does not require pre-**

			shipment inspection • **CLOTRIMAZOLE 2% VAGINAL CREAM 30GM** **This item does not require pre-shipment inspection** • **ENALAPRIL 5MG TABLETS** **This item does not require pre-shipment inspection** • **ALBENDAZOLE SUSPENSION 100MG/5ML IN 10ML BOTTLE** **This item does not require pre-shipment inspection** • **CIPROFLOXACIN 200MG/100ML INJECTION** **This item does not require pre-shipment inspection** • **METOCLOPRAMIDE 10MG TABLETS** **This item does not require pre-shipment inspection** • **TIMOLOL 0.5% EYE DROPS, 5MLS** **This item does not require pre-shipment inspection** • **DANAZOL 100MG TABLET** **This item does not require pre-shipment inspection** • **ANTI-HAEMORRHOIDS SUPPOSITORIES** **This item does not require pre-shipment inspection**

			• **BENZATHINE PENICILLIN FORTIFIED PDR F INJ 2.4 MU** **This item does not require pre-shipment inspection** • **AMPICILLIN + CLOXACILLIN 125MG/5ML SUSPENSION** **This item does not require pre-shipment inspection** • **LIDOCAINE 5%+DEXTROSE 7.5 % SPINAL INJECTION.** **This item does not require pre-shipment inspection** • **WATER FOR INJECTION 10ML** **This item does not require pre-shipment inspection** • **PHYTOMENADIONE(VITAMIN K1) INJECTION** **This item does not require pre-shipment inspection** • **ALUMINIUM HYDROXIDE355MG + MAGNESIUM HYDROXIDE 60MG + SIMETHICONE 100MG + LIQUORICE 400MG (RELCER GEL) 180MLS** **This item does not require pre-shipment inspection** • **SILVER SULFADIAZINE 1% + CHLORHEXIDINE 20GM CREAM** **This item does not require pre-shipment inspection** • **HYOSCINE -N- BUTYLBROMIDE**

			10MG TABS **This item does not require pre-shipment inspection** • **CLOTRIMAZOLE CREAM 1% (PROCOSTAT)** **This item does not require pre-shipment inspection** • **HYDROCORTISONE 1%, 15 GRAMS, CREAM** **This item does not require pre-shipment inspection** • **ATROPINE 1MG/ML, 1ML, INJECTION.** **This item does not require pre-shipment inspection** • **FLUCYTOSINE 250MG TABLET** **This item does not require pre-shipment inspection** • **LIVOLIN FORTE (MULTIVITAMIN) CAPSULE** **This item does not require pre-shipment inspection** • **BACLOFEN 10MG TABLETS** **This item does not require pre-shipment inspection** • **HALOTHANE INHALATION** **This item does not require pre-shipment inspection** • **HYDROCORTISONE 0.01%,15G CREAM**

			This item does not require pre-shipment inspection • **MIDAZOLAM 5MG/ML INJECTION** **This item does not require pre-shipment inspection** • **MICONAZOLE NITRATE 400MG PESSARIES (GYNAZOL)** **This item does not require pre-shipment inspection** • **METFORMIN 850MG TABLET** **This item does not require pre-shipment inspection** • **PREGABALIN 75MG CAPSULE** **This item does not require pre-shipment inspection** • **LACTULOSE 3.4GM/5ML SOLUTION** **This item does not require pre-shipment inspection** • **CARBAMAZAPINE 200MG TABLETS** **This item does not require pre-shipment inspection** • **MOMETASONE NASAL SPRAY** **This item does not require pre-shipment inspection** • **DOXYLAMINE SUCCINATE 10MG + PYRIDOXINE HCL 10MG TABLETS** **This item does not require pre-shipment inspection** • **DIAZEPAM 5MG TABLETS**

			This item does not require pre-shipment inspection • **ISOSORBIDE DINITRATE 10MG TABLETS** **This item does not require pre-shipment inspection** • **FLUCLOXACILLIN 250MG CAPSULE** **This item does not require pre-shipment inspection** • **CLONIDINE 500µG/ML INJECTION.** **This item does not require pre-shipment inspection** • **DICLOFENAC SODIUM 50MG TABLET** **This item does not require pre-shipment inspection** • **CLOPIDOGREL 75MG TABLETS** **This item does not require pre-shipment inspection** • **BETAMETHASONE 0.1 + NEOMYCIN 0.5% CREAM** **This item does not require pre-shipment inspection** • **RABIES VACCINE USP (POTENCY OF RABIES ANTIGEN ≥ 2.5 IU/DOSE** **This item does not require pre-shipment inspection** • **ENALAPRIL 2.5MG TABLET**

			This item does not require pre-shipment inspection • **EPLERENONE 50 MG TABLET** **This item does not require pre-shipment inspection** • **BETAMETHASONE 0.1%, 15GM CREAM** **This item does not require pre-shipment inspection** • **GENTIAN VIOLET 30ML** **This item does not require pre-shipment inspection** • **CARBAMAZEPINE 100MG TABLET** **This item does not require pre-shipment inspection** • **KETOPROFEN 50MG CAPSULES** **This item does not require pre-shipment inspection** • **CALCIUM + VITAMIN D3+VITAMIN A CAPSULE** **This item does not require pre-shipment inspection** • **VITAMIN B COMPLEX TABLET.** **This item does not require pre-shipment inspection** • **IPRATROPIUM NEBULIZER SOLUTION 500MCG/2ML (IPRAVENT)** **This item does not require pre-shipment inspection**

			• **VITAMIN B COMPLEX SYRUP, 100ML** **This item does not require pre-shipment inspection** • **BUDESONIDE INHALER 100MCG PER DOSE (200 DOSES)** **This item does not require pre-shipment inspection** • **ALUMINIUM HYDROXIDE + MAGNESIUM HYDROXIDE SUSPENSION, 100MLS** **This item does not require pre-shipment inspection** • **ORAL REHYDRATION SALTS (ORS) FOR 1 LITRE POWDER** **This item does not require pre-shipment inspection** • **TELMISARTAN 40MG TABLET** **This item does not require pre-shipment inspection** • **MEFENAMIC ACID 500MG TABLET** **This item does not require pre-shipment inspection** • **MANNITOL 20% 100MLS SOLUTION FOR IV INFUSION (ABTOL)** **This item does not require pre-shipment inspection** • **LABETALOL 10MG/ML INJECTION** **This item does not require pre-shipment inspection**

			• **GLIMEPIRIDE 1+ METFORMIN 500MG TABLETS (GEMER 1)** **This item does not require pre-shipment inspection** • **PRAZIQUANTEL 600MG TABLET** **This item does not require pre-shipment inspection** • **CANDERSARTAN 8MG TABLET** **This item does not require pre-shipment inspection** • **METRONIDAZOLE 200MG/5ML 100ML SUSPENSION** **This item does not require pre-shipment inspection** • **LORAZEPAM 2MG/ML IN 1ML AMPOULE INJECTION** **This item does not require pre-shipment inspection** • **BENZYL PENICILLIN PDR F 5 MU INJECTION** **This item does not require pre-shipment inspection** • **CARVEDILOL 6.25MG TABLET** **This item does not require pre-shipment inspection** • **ALBENDAZOLE 200MG TABLETS** **This item does not require pre-shipment inspection** • **ARTEMETHER/LUMEFANTRINE 20/120 MG TABLETS**

			This item does not require pre-shipment inspection • **LOW MOLECULAR WEIGHT HEPARIN 25MG INJECTION** **This item does not require pre-shipment inspection** • **GLICLAZIDE 40MG TABLET** **This item does not require pre-shipment inspection** • **METRONIDAZOLE 1% GEL (CORNIZOLE)** **This item does not require pre-shipment inspection** • **HYDROGEN PEROXIDE MOUTHWASH 3% SOLUTION, 100MLS** **This item does not require pre-shipment inspection** • **DEXTROSE 5% (D5) INTRAVENOUS INFUSION 500MLS** **This item does not require pre-shipment inspection** • **SALBUTAMOL 4MG TABLETS** **This item does not require pre-shipment inspection** • **ATENOLOL 50MG TABLETS** **This item does not require pre-shipment inspection** • **TERBINAFINE 250MG TABLET.** **This item does not require pre-**

			shipment inspection • **BENZOYL PEROXIDE 2.5% ,20GM GEL** **This item does not require pre-shipment inspection** • **PROMETHAZINE 25MG TABLETS** **This item does not require pre-shipment inspection** • **PARACETAMOL 125MG SUPPOSITORIES** **This item does not require pre-shipment inspection** • **CLARYTHROMYCIN 500MG + TINIDAZOLE 500MG + LANSOPRAZOLE 30MG (42 TABLETS** **This item does not require pre-shipment inspection** • **HYDROGEN PEROXIDE 3%, 10MLS EAR DROPS SOLUTION** **This item does not require pre-shipment inspection** • **ESOMEPRAZOLE 20MG TABLET** **This item does not require pre-shipment inspection** • **CEFIXIME 400MG TABLETS** **This item does not require pre-shipment inspection** • **MONTELUKAST 10MG TABLETS** **This item does not require pre-**

			shipment inspection • **BISACODYL 5MG TABLET** **This item does not require pre-shipment inspection** • **BUPIVACAINE 0.5% (HEAVY MARCAINE) HYPERBARIC, 4ML INJECTION** **This item does not require pre-shipment inspection** • **LEVOTHYROXINE 25MCG TABLET** **This item does not require pre-shipment inspection** • **BISOPROLOL FUMARATE 10 MG TABLET** **This item does not require pre-shipment inspection** • **CALCIUM GLUCONATE 100MG/ML 10ML INJECTION** **This item does not require pre-shipment inspection** • **CEFIXIME 200MG TABLET** **This item does not require pre-shipment inspection** • **POTASSIUM PERMANGANATE 1:4000,100ML SOLUTION** **This item does not require pre-shipment inspection** • **CHLORAMPHENICOL 5% EAR/EYE DROPS** **This item does not require pre-**

			shipment inspection • **BENZOIC ACID COMPOUND (WHITFIELD) OINTMENT 6%+3%** **This item does not require pre-shipment inspection** • **GENTAMYCIN 40MG/ML, 2ML INJECTION** **This item does not require pre-shipment inspection** • **DESLORATIDINE 5MG TABLETS** **This item does not require pre-shipment inspection** • **PODOPHYLINE TOPICAL SOLUTION 10-25%** **This item does not require pre-shipment inspection** • **ASCORBIC ACID 500MG TABLET** **This item does not require pre-shipment inspection** • **CLOTRIMAZOLE EAR DROPS 15MLS** **This item does not require pre-shipment inspection** • **BRIMONIDINE TARTATE 0.2% EYE DROPS** **This item does not require pre-shipment inspection** • **ECONAZOLE 5% EYE DROP** **This item does not require pre-shipment inspection**

			• **ACYCLOVIR 10G CREAM** **This item does not require pre-shipment inspection** • **CEPHALEXIN 250MG CAPSULES** **This item does not require pre-shipment inspection** • **FLUCONAZOLE 150MG TABLETS** **This item does not require pre-shipment inspection** • **DIMERCAPSULEROL 50MG/ML 2ML INJECTION** **This item does not require pre-shipment inspection** • **PHENYTOIN 100MG TABLETS** **This item does not require pre-shipment inspection** • **DIHYDROGESTERONE 10MG TABLETS** **This item does not require pre-shipment inspection** • **BUPIVACAINE (LIGHT MARCAINE) 5MG/5ML INJECTION** **This item does not require pre-shipment inspection** • **BUDESONIDE INHALER 200MCG PER DOSE (200 DOSES)** **This item does not require pre-shipment inspection** • **MICONAZOLE CREAM 2%, 20GM** **This item does not require pre-**

			shipment inspection • **OXYTOCIN 10IU/ML, 1ML INJECTION** **This item does not require pre-shipment inspection** • **ANTI RHO-D HUMAN IMMUNOGLOBIN MONOCLONAL 150MCG INJECTION** **This item does not require pre-shipment inspection** • **DONEPEZIL 5MG TABLET** **This item does not require pre-shipment inspection** • **FUROSEMIDE 40MG TABLETS** **This item does not require pre-shipment inspection** • **CEFTAZIDIME SODIUM 250MG INJECTION** **This item does not require pre-shipment inspection** • **IMIPRAMINE 25MG TABLET** **This item does not require pre-shipment inspection** • **CLOPIDOGREL 300MG TABLET** **This item does not require pre-shipment inspection** • **ATROPINE 1% EYE DROPS** **This item does not require pre-shipment inspection**

			• **EPHEDRINE 30MG/ML, 1ML INJECTION** **This item does not require pre-shipment inspection** • **BROMOCRIPTINE MESYLATE 2.5MG TABLET** **This item does not require pre-shipment inspection** • **ARTEMETHER/LUMEFANTRINE 20/120 MG TABLETS** **This item does not require pre-shipment inspection** • **DEXTROSE 50%, 250ML SOLUTION FOR INTRAVENOUS INJECTION/INFUSION** **This item does not require pre-shipment inspection** • **FLUOXETINE 20MG CAPSULE** **This item does not require pre-shipment inspection** • **TIMOLOL 0.25% EYE DROPS, 10MLS** **This item does not require pre-shipment inspection** • **DICLOFENAC SODIUM 25MG/ML, 3ML INJECTION** **This item does not require pre-shipment inspection** • **SALBUTAMOL 2.5MG NEBULIZER** **This item does not require pre-shipment inspection**

			• **GLICLAZIDE 60MG TABLETS** **This item does not require pre-shipment inspection** • **ISOSORBIDE MONONITRATE 10MG TABLET** **This item does not require pre-shipment inspection** • **PHENOBARBITAL SODIUM 100MG/ML 2ML INJECTION** **This item does not require pre-shipment inspection** • **CO-PACK FOR DIARRHOEA TREATMENT KIT (ORS+ZINC 20MG) TABLET** **This item does not require pre-shipment inspection** • **DICLOFENAC GEL 30G, 1% W/W** **This item does not require pre-shipment inspection** • **AMPICILLIN + CLOXACILLIN POWDER FOR INJECTION 500MG** **This item does not require pre-shipment inspection** • **GLIBENCLAMIDE 5MG TABLETS** **This item does not require pre-shipment inspection** • **DOBUTAMINE 12.5 MGS/ML INJECTION 20 MLS.** **This item does not require pre-shipment inspection**

			• **HYDROXYCOBALAMINE(VITAMIN B 12) 1000 MCG,INJECTION.** **This item does not require pre-shipment inspection** • **SILVER SULPHADIAZINE 1% W/W CREAM** **This item does not require pre-shipment inspection** • **CHLORAMPHENICOL + DEXAMETHASONE EYE DROPS, 5MLS** **This item does not require pre-shipment inspection** • **LIGNOCAINE HYDROCHLORIDE INJECTION 1% 10ML** **This item does not require pre-shipment inspection** • **CHLORPHENIRAMINE 10MG/ML INJECTION, 1ML** **This item does not require pre-shipment inspection** • **CHLORPROMAZINE 25MG TABLET** **This item does not require pre-shipment inspection** • **BENZOYL PEROXIDE 10% CREAM, 20GM** **This item does not require pre-shipment inspection** • **METHYLDOPA 250MG TABLETS** **This item does not require pre-**

			shipment inspection • **ACTIVATED CHARCOAL 200 MG (WELLVITA)** **This item does not require pre-shipment inspection** • **DEXAMETHASONE 0.1% + CHLORAMPHENICOL EYE DROP 1%W/V 5ML** **This item does not require pre-shipment inspection** • **CARBIMAZOLE 5MG TABLETS** **This item does not require pre-shipment inspection** • **CETIRIZINE HYDROCHLORIDE 5MG/5ML ORAL LIQUID** **This item does not require pre-shipment inspection** • **2,3-DIMERCAPSULETO-SUCCINIC ACID 100MG CAPSULES** **This item does not require pre-shipment inspection** • **LORAZEPAM 2 MG TABLETS (LORIVAN 2)** **This item does not require pre-shipment inspection** • **ONDANSETRON 8MG TABLET** **This item does not require pre-shipment inspection** • **DILTIAZEM 60MG TABLETS.** **This item does not require pre-**

			shipment inspection • **SPIRONOLACTONE 25MG TABLETS** **This item does not require pre-shipment inspection** • **HYDROGEN PEROXIDE 6% 100ML** **This item does not require pre-shipment inspection** • **FINASTERIDE 5MG TABLETS** **This item does not require pre-shipment inspection** • **TELMISARTAN 80MG TABLET** **This item does not require pre-shipment inspection** • **ERYTHROMYCIN 125MG/5ML 100ML** **This item does not require pre-shipment inspection** • **INSULIN HUMAN ZINC SUSPENSION INJECTION 100IU (LENTE)** **This item does not require pre-shipment inspection** • **MEBENDAZOLE 100MG TABLETS** **This item does not require pre-shipment inspection** • **ACYCLOVIR 200MG TABLETS** **This item does not require pre-shipment inspection** • **AMOXYCILLIN + CLAVULANIC**

			ACID INJECTION 1.2GM INJECTION **This item does not require pre-shipment inspection** • **NIFEDIPINE 10MG SLOW RELEASE TABLET** **This item does not require pre-shipment inspection** • **DEXAMETHASONE 0.5MG TABLET** **This item does not require pre-shipment inspection** • **MANNITOL 20% 500MLS INJECTION** **This item does not require pre-shipment inspection** • **CYCLOPENTOLATE 1% EYE DROP** **This item does not require pre-shipment inspection** • **CALCIUM 500MG+ VITAMIN D 10MCG TABLET** **This item does not require pre-shipment inspection** • **HEPARIN 5000IU/ML, 5MLS INJECTION** **This item does not require pre-shipment inspection** • **MICONAZOLE 2% ORAL GEL 30G** **This item does not require pre-shipment inspection** • **LIGNOCAINE 2% , 50ML**

			INJECTION **This item does not require pre-shipment inspection** • **LANSOPRAZOLE 30MG TABLET** **This item does not require pre-shipment inspection** • **CANDESARTAN 16MG + HYDROCHLOROTHIAZIDE 12.5MG TABLET** **This item does not require pre-shipment inspection** • **IPRATROPIUM BROMIDE AEROSOL 20MCG INHALATION (AEROSOL)** **This item does not require pre-shipment inspection** • **BENDROFLUAZIDE 5MG TABLETS** **This item does not require pre-shipment inspection** • **COUGH EXPECTORANT CHILD SYRUP100ML** **This item does not require pre-shipment inspection** • **PANTOPRAZOLE 40MG INJECTION (PEPRASOL)** **This item does not require pre-shipment inspection** • **DEXTROSE NORMAL SALINE (DNS) INJECTION 500 MLS** **This item does not require pre-shipment inspection**

			• **AMLODIPINE 5MG TABLETS** **This item does not require pre-shipment inspection** • **COUGH EXPECTORANT ADULT SYRUP,100MLS** **This item does not require pre-shipment inspection** • **SODIUM VALPROATE 200MG TABLETS** **This item does not require pre-shipment inspection** • **TRAMADOL 50MG/ML, 2ML INJECTION** **This item does not require pre-shipment inspection** • **METOCLOPRAMIDE 5MG/5ML 100ML SYRUP** **This item does not require pre-shipment inspection** • **PANCURONIUM BROMIDE 2MG/ML 2ML INJECTION** **This item does not require pre-shipment inspection** • **PROPRANOLOL 40MG TABLET** **This item does not require pre-shipment inspection** • **NORFLOXACIN 400MG TABLET** **This item does not require pre-shipment inspection** • **EPHEDRINE NASAL DROPS 0.5%**

			(PAEDIATRIC) **This item does not require pre-shipment inspection** • **DICYCLOMINE HCL 20MG + PARACETAMOL 500MG TABLETS** **This item does not require pre-shipment inspection** • **BENZHEXOL 5MG TABLETS** **This item does not require pre-shipment inspection** • **TRIAMCINOLONE INJECTION 40MG/ML (TRICORT 40)** **This item does not require pre-shipment inspection** • **DOXYCYCLINE 100MG CAPSULES** **This item does not require pre-shipment inspection** • **SODIUM CHLORIDE 0.9% 500ML SOLUTION FOR INTRAVENOUS INFUSSION, 500ML** **This item does not require pre-shipment inspection** • **PHENOBARBITAL 100MG TABLETS** **This item does not require pre-shipment inspection** • **LINDANE LOTION 100ML** **This item does not require pre-shipment inspection** • **MONTELUKAST 5MG TABLETS** **This item does not require pre-**

			shipment inspection • **CALCIUM GLUCONATE 500MG TABLET** **This item does not require pre-shipment inspection** • **ISOSORBIDE DINITRATE 20MG TABLET** **This item does not require pre-shipment inspection** • **CARVEDILOL 12.5MG TABLETS** **This item does not require pre-shipment inspection** • **ZINC SULPHATE 20MG TABLETS** **This item does not require pre-shipment inspection** • **FERROUS SULPHATE 200MG + FOLIC ACID 0.25 MG TABLETS** **This item does not require pre-shipment inspection** • **HYDRALAZINE 25MG TABLETS** **This item does not require pre-shipment inspection** • **CALCIUM CARBONATE 500MG TABLETS** **This item does not require pre-shipment inspection** • **CHLORPROMAZINE 25MG/ML 2ML INJECTION** **This item does not require pre-shipment inspection**

			• **CALCIUM + MAGNESIUM + ZINC + VITAMIN D3 TABLET** **This item does not require pre-shipment inspection** • **BISOPROLOL FUMARATE 5MG TABLETS** **This item does not require pre-shipment inspection** • **METRONIDAZOLE 5MG/ML, 100ML SOLUTION FOR INTRAVENOUS INJECTION/INFUSION** **This item does not require pre-shipment inspection** • **DICLOFENAC SODIUM 100MG SLOW RELEASE** **This item does not require pre-shipment inspection** • **PROMETHAZINE HCL 25MG/ML, 2ML INJECTION** **This item does not require pre-shipment inspection** • **FLUMAZENIL INJ** **This item does not require pre-shipment inspection** • **BENZYL BENZOATE EMULSION (BBE) 25%W/V** **This item does not require pre-shipment inspection** • **CYCLOPENTOLATE 0.5% EYE DROP** **This item does not require pre-**

			shipment inspection • **MELOXICAM 15MG TABLETS** **This item does not require pre-shipment inspection** • **CLOTRIMAZOLE 1%W/W, 30GM POWDER** **This item does not require pre-shipment inspection** • **CLINDAMYCIN IV 150MG/ML INJECTION,2MLS** **This item does not require pre-shipment inspection** • **FRUSEMIDE 10MG/ML, 2ML INJECTION** **This item does not require pre-shipment inspection** • **CLARITHROMYCIN 500MG TABLET** **This item does not require pre-shipment inspection** • **PROPOFOL 1% MG/ML INJECTION** **This item does not require pre-shipment inspection** • **DORZOLAMIDE 2% EYE DROPS (OCUDOR)** **This item does not require pre-shipment inspection** • **DOXAZOCIN 2MG TABLET** **This item does not require pre-shipment inspection**

			• **ACETAZOLAMIDE 250MG TABLET** **This item does not require pre-shipment inspection** • **HALOPERIDOL 1.5MG TABLETS** **This item does not require pre-shipment inspection** • **DEXTROSE SOLUTION 10%,500ML INTRAVENOUS INFUSION** **This item does not require pre-shipment inspection** • **SODIUM DICHLOROISOCYANURATE 2.7GM TABLETS** **This item does not require pre-shipment inspection** • **HYOSCINE BUTYLBROMIDE 20MG/ML, 1ML AMPOULE** **This item does not require pre-shipment inspection** • **POVIDONE IODINE LIQUID 10%, 250ML** **This item does not require pre-shipment inspection** • **ANTITHYMOCITE GLOBULIN 50MG/ML, 5ML (ATG) INJECTION** **This item does not require pre-shipment inspection** • **KETOPROFEN 30GM GEL (2.5% w/w)** **This item does not require pre-shipment inspection**

			• **NITROFURANTOIN 100MG TABLETS** **This item does not require pre-shipment inspection** • **GRISEOFULVIN 500MG TABLETS** **This item does not require pre-shipment inspection** • **FLUCONAZOLE 200MG TABLET** **This item does not require pre-shipment inspection** • **CIPROFLOXACIN 0.3% EYE DROPS 10MLS** **This item does not require pre-shipment inspection** • **IRBESARTAN 300MG TABLET** **This item does not require pre-shipment inspection** • **BETAHISTIDINE 8 MG (BETASERC)TABS** **This item does not require pre-shipment inspection** • **GENTAMYCIN 2ML INJ 10MG/ML** **This item does not require pre-shipment inspection** • **GLUTARALDEHYDE WITH ACTIVATOR 2.45% w/v, 5L SOLUTION** **This item does not require pre-shipment inspection** • **GLIMEPIRIDE 2MG TABLET**

			This item does not require pre-shipment inspection • **LORAZEPAM 1MG TABS** **This item does not require pre-shipment inspection** • **ATROPINE 0.5% EYE DROP** **This item does not require pre-shipment inspection** • **PREDNISOLONE 0.5% EYE DROPS** **This item does not require pre-shipment inspection** • **CARBAMAZEPINE SYRUP 100MG/5ML, 100ML** **This item does not require pre-shipment inspection** • **MULTIVITAMIN SYRUP 100ML** **This item does not require pre-shipment inspection** • **AMOXICILLIN 250MG CAPSULES** **This item does not require pre-shipment inspection** • **ERGOCALCIFEROL (VITAMIN D) 1.25MG (50000IU) CAPSULE** **This item does not require pre-shipment inspection** • **ESMOLOL 10MG/ML.10ML** **This item does not require pre-shipment inspection** • **CLONIDINE 100MCG TABLET**

			This item does not require pre-shipment inspection • **SNAKE VENOM POLYVALENT ANTISERUM 10MLS, VIAL.** **This item does not require pre-shipment inspection** • **CHLORPHENIRAMINE MALEATE SYRUP 2MG/5ML, 100ML** **This item does not require pre-shipment inspection** • **NORETHISTERONE 5MG TABLET** **This item does not require pre-shipment inspection** • **ACETYLSALICYLIC ACID (ENTERIC COATED) 75MG TABLETS** **This item does not require pre-shipment inspection** • **PARACETAMOL SYRUP 120MG/5MLS,100MLS** **This item does not require pre-shipment inspection** • **MAGNESIUM TRISILICATE BP COMPOUND TABLETS** **This item does not require pre-shipment inspection** • **LOSARTAN 50MG TABLET** **This item does not require pre-shipment inspection** • **ALLOPURINOL 100MG TABLET**

			This item does not require pre-shipment inspection • **AMLODIPINE 10MG TABLET** **This item does not require pre-shipment inspection** • **CLONAZEPAM 0.5MG TABLETS** **This item does not require pre-shipment inspection** • **CEFIPIME 1GM INJECTION** **This item does not require pre-shipment inspection** • **FOLIC ACID 5MG TABLETS** **This item does not require pre-shipment inspection** • **METOCLOPRAMIDE 5MG/ML ,2ML INJECTION** **This item does not require pre-shipment inspection** • **TRETINOIN 0.05% CREAM** **This item does not require pre-shipment inspection** • **LORATADINE 10MG TABLETS (LORFAST)** **This item does not require pre-shipment inspection** • **LEVODOPA 100 + CARBIDOPA 25MG TABLETS** **This item does not require pre-shipment inspection**

			• **PHENOXYMETHYLPENICILLIN 250MG TABLETS** **This item does not require pre-shipment inspection** • **SUXAMETHONIUM CHLORIDE 50MG/ML INJECTION, 2ML** **This item does not require pre-shipment inspection** • **N-ACETYLCYSTEINE 200MG/ML, 10ML INJECTION** **This item does not require pre-shipment inspection** • **BETAMETHASONE 4MG/ML INJECTION** **This item does not require pre-shipment inspection** • **PANTOPRAZOLE 40MG TABLETS** **This item does not require pre-shipment inspection** • **CLINDAMYCIN 150MG CAPSULE** **This item does not require pre-shipment inspection** • **TELMISARTAN 40 MG + HYDROCHLOROTHIAZIDE 12.5 MG TABLET** **This item does not require pre-shipment inspection** • **CHLORAMPHENICOL 1% EYE DROP** **This item does not require pre-shipment inspection**

			• **CIPROFLOXACIN 500MG TABLETS** **This item does not require pre-shipment inspection** • **LORATADINE 5MG/5ML SYRUP (LORATA)** **This item does not require pre-shipment inspection** • **DOMPERIDONE 10MG TABLETS** **This item does not require pre-shipment inspection** • **ADRENALINE 1MG/ML, 1ML INJECTION** **This item does not require pre-shipment inspection** • **IBUPROFEN 100MG/5ML SYRUP 100ML** **This item does not require pre-shipment inspection** • **AMOXICILLIN 500MG + CLAVULANIC ACID 125MG (AUGUMENTIN 625) TABLETS** **This item does not require pre-shipment inspection** • **PREDNISOLONE 5MG TABLETS** **This item does not require pre-shipment inspection** • **AMITRYPTILINE 25MG TABLETS** **This item does not require pre-shipment inspection** • **LAMOTRIGINE 100MG TABLETS**

			This item does not require pre-shipment inspection • **EPHEDRINE NASAL DROPS 1% (ADULT), 15MLS** **This item does not require pre-shipment inspection** • **IBUPROFEN 400MG CAPSULES (IBUBERG)** **This item does not require pre-shipment inspection** • **DEXMEDETOMIDINE 200MG/2ML INJECTION** **This item does not require pre-shipment inspection** • **CIPROFLOXACIN EAR DROPS** **This item does not require pre-shipment inspection** • **AMIODARONE 50MG/ML, 3MLS INJECTION** **This item does not require pre-shipment inspection** • **Flucytocine 500mg Tablet** **This item does not require pre-shipment inspection** • **MEFENAMIC ACID 250MG CAPSULES** **This item does not require pre-shipment inspection** • **DIAZEPAM 5 MG/ML, 2ML INJECTION**

			This item does not require pre-shipment inspection • **CALCIUM +VITAMIN B+ VITAMIN C TABLET** **This item does not require pre-shipment inspection** • **TETRACYCLINE EYE OINTMENT 1% 3.5GM** **This item does not require pre-shipment inspection** • **CEFTRIAXONE 1G POWDER FOR INJECTION** **This item does not require pre-shipment inspection** • **PHENOXYMETHYLPENICILLIN POWDER FOR ORAL SOLUTION 125MG/5ML** **This item does not require pre-shipment inspection** • **RABIES VACCINE USP (POTENCY OF RABIES ANTIGEN ≥ 2.5 IU/DOSE** **This item does not require pre-shipment inspection** • **CHLOROXYLENOL (DETTOL) LIQUID 5%** **This item does not require pre-shipment inspection** • **BETAXOLOL 0.25% - 0.5% EYE DROP**

			This item does not require pre-shipment inspection • **METOPROL 50 MG TABLETS (METAPROL)** **This item does not require pre-shipment inspection** • **FLUTICASONE FUORATE NASAL SPRAY** **This item does not require pre-shipment inspection** • **KETOPROFEN 75MG CAPSULES** **This item does not require pre-shipment inspection** • **TACROLIMUS 0.1% OINTMENT** **This item does not require pre-shipment inspection** • **CALCIUM CHLORIDE INJECTION 10%, 10MLS** **This item does not require pre-shipment inspection** • **ESOMEPRAZOLE 20MG TABLET** **This item does not require pre-shipment inspection** • **LIGNOCAINE HEAVY SPINAL 5%** **This item does not require pre-shipment inspection** • **XYLOMETAZOLINE 0.1%,10 MLS NASAL SPRAY (XYLOACINO)** **This item does not require pre-shipment inspection**

			• **ATORVASTATIN 20MG TABLETS** **This item does not require pre-shipment inspection** • **ALBENDAZOLE 400MG TABLETS** **This item does not require pre-shipment inspection** • **FORMALDEHYDE 36-37% LIQUID** **This item does not require pre-shipment inspection** • **ITRACONAZOLE 100 MG CAPSULE** **This item does not require pre-shipment inspection** • **NEOSTIGMINE 2.5MG/ML, 1ML INJECTION** **This item does not require pre-shipment inspection** • **LEVETIRACETAM 250MG TABLET** **This item does not require pre-shipment inspection** • **AMOXICILLIN + CLAVULIN ACID (228MG/5ML) 100ML, SYRUP** **This item does not require pre-shipment inspection** • **ATROPINE 0.5MG/ML, 1ML, INJECTION.** **This item does not require pre-shipment inspection** • **ERGOCALCIFEROL (VITAMIN D) 0.25MG/ML (10000IU/ML) ORAL SOLUTION**

			This item does not require pre-shipment inspection • **PARACETAMOL 500MG TABLETS** **This item does not require pre-shipment inspection** • **DICYCLOMINE 20MG INJECTION** **This item does not require pre-shipment inspection** • **THIOPENTAL SODIUM 500MG, PDR F INJ** **This item does not require pre-shipment inspection** • **LISINOPRIL 10MG TABLET** **This item does not require pre-shipment inspection** • **FEBUXOSTAT 80MG TABLET** **This item does not require pre-shipment inspection** • **NYSTATIN SUSPENSION 100000 IU/ML 30ML** **This item does not require pre-shipment inspection** • **LOSARTAN 50MG + HYDROCHLORTHIAZIDE 12.5MG TABLETS** **This item does not require pre-shipment inspection** • **ATRACURIUM 50MG/5ML, 5ML INJECTION** **This item does not require pre-**

			shipment inspection • **EPLERENONE 25 MGS, TABLET** **This item does not require pre-shipment inspection** • **DICLOFENAC POTASSIUM 50MG TABLET** **This item does not require pre-shipment inspection** • **DEXAMETHASONE 0.1% + NEOMYCIN 0.35% EAR/EYE DROPS (DEXANEO), 10MLS** **This item does not require pre-shipment inspection** • **BETAMETHASONE + GENTAMYCIN EYE DROPS** **This item does not require pre-shipment inspection** • **OLANZAPINE 10MG TABLETS** **This item does not require pre-shipment inspection** • **CHLORAMPHENICOL 0.5% EYE OINTMENT** **This item does not require pre-shipment inspection** • **CALCIUM +VITAMIN A CAPSULE** **This item does not require pre-shipment inspection** • **HEPARIN GEL** **This item does not require pre-shipment inspection**

			• **COAL TAR 5% TOPICAL SOLUTION** **This item does not require pre-shipment inspection** • **CO-TRIMOXAZOLE SUSPENSION 200/40MG/5ML, 100MLS** **This item does not require pre-shipment inspection** • **ATORVASTATIN 10MG TABLET** **This item does not require pre-shipment inspection** • **POWDER WASHING MACHINE SOAP (SILEX)** **This item does not require pre-shipment inspection** • **GLIMEPIRIDE 1MG TABLET** **This item does not require pre-shipment inspection** • **IRBESARTAN 150MG TABLETS (IROVEL)** **This item does not require pre-shipment inspection** • **DEFEROXAMINE 500MG POWDER FOR (MESYLATE) INJECTION** **This item does not require pre-shipment inspection** • **TRANEXAMIC ACID 100MG/ML 5ML INJECTION** **This item does not require pre-shipment inspection** • **ARTEMETHER/LUMEFANTRINE**

			20/120 MG TABLETS **This item does not require pre-shipment inspection** • **VITAMIN B COMPLEX 2ML INJECTION** **This item does not require pre-shipment inspection** • **AMIODARONE 100MG TABLET** **This item does not require pre-shipment inspection** • **GENTAMYCIN 20MG/ML, 2ML INJECTION** **This item does not require pre-shipment inspection** • **ADENOSINE 3MG/ML IN SALINE INJECTION** **This item does not require pre-shipment inspection** • **HAND SANITIZER ANTISEPTIC, 250ML** **This item does not require pre-shipment inspection** • **TINIDAZOLE 500MG TABLET** **This item does not require pre-shipment inspection** • **MEROPENEM TRIHYDRATE 1GM INJECTION (POWDER FOR RECONSTITUTION)** **This item does not require pre-shipment inspection**

			• **AZITHROMYCIN SUSPENSION 200MG/5MLS 15ML** **This item does not require pre-shipment inspection** • **URSODEOXYCHOLIC ACID 300MG TABLET** **This item does not require pre-shipment inspection** • **DUTASTERIDE 0.5MG CAPSULE** **This item does not require pre-shipment inspection** • **HYDROCHLOROTHIAZIDE 25MG TABLETS** **This item does not require pre-shipment inspection** **NONE**
15.	Packing Requirements	12.2	NONE
16.	Delivery and Documents	13.1 & 13.3	Sample provision (CIF/CIP terms) For Goods supplied form abroad: Upon shipment, the Supplier shall notify the Purchaser and the insurance company in writing the full details of the shipment including Contract number, description of the Goods, quantity, date and place of shipment, mode of transportation, and estimated date of arrival at place of destination. In the event of Goods sent by airfreight, the Supplier shall notify the Purchaser a minimum of forty-eight (48) hours ahead of dispatch, the name of the

			carrier, the flight number, the expected time of arrival, and the waybill number. The Supplier shall fax and them send by courier the following documents to the Purchaser, with a copy to the insurance company: i) three originals and two copies of the Supplier's invoice, showing Purchaser as [enter correct description of Purchaser for customs purposes]; the Contract number, loan number, Goods description, quantity, unit price, and total amount. Invoices must be signed in original, stamped, or sealed with the company stamp/seal. ii) one original and two copies of the negotiable, clean, on-board through bill of lading marked "freight prepaid" and showing Purchaser as [enter correct number of Purchaser for customs purposes] and Notify Party as stated in the Contract, with delivery through to final destination as per the Schedule of Requirements and two copies of non-negotiable bill of lading, or three copies of railway consignment note, road consignment note, truck or air waybill, or multimodal transport document, marked "freight prepaid" and showing delivery through to final destination as per the Schedule of Requirements; iii) four copies of the packing list identifying contents of each package; iv) copy of the Insurance Certificate, showing the Purchaser as the beneficiary; v) one original of the manufacturer's or Supplier's Warranty Certificate covering

			all items supplied; vi) one original of the Supplier's Certificate of Origin covering all items supplied; vii) original copy of the Certificate of Inspection furnished to Supplier by the nominated inspection agency and six copies (where inspection is required); For goods shipped by sea the following documents to be submitted immediately by email and original documents to be submitted at least 14 days prior to ship arrival. viii) Certificate of analysis/ conformity for medical supplies and equipment to be included as the delivery document ix) EFD receipt to be included for VAT registered suppliers (local) x) Endorsed Import permit to be included in the delivery documents for imported goods. xi) any other procurement-specific documents required for delivery / payment purposes. For Goods from within the United Republic of Tanzania. Upon or before delivery of the Goods, the Supplier shall notify the Purchaser in writing and deliver the following documents to the Purchaser: (i) two originals and two copies of the Supplier's invoice, showing Purchaser, the Contract number, loan number; Goods' description, quantity, unit price, and total

			amount. Invoices must be signed in original and stamped or sealed with the company stamp / seal; ii) two copies of delivery note, railway consignment note, road consignment note, truck or air waybill, or multimodal transport document showing Purchaser as [enter correct name of Purchaser for customs purposes] and delivery through to final destination as stated in the Contract. iii) copy of the Insurance Certificate, showing the Purchaser as the beneficiary, iv) four copies of the packing list identifying contents of each package; v) one original of the manufacturer's or Supplier's Warranty certificate covering all items supplied; vi) one original of the Supplier's Certificate of Origin covering all items supplied' vii) original copy of the Certificate of Inspection furnished to Supplier by the nominated inspection agency and six copies (where inspection is required) viii) other procurement-specific documents required for delivery / payment purposes - Not Applicable
17.	Insurance	14.1	TZS 0.00
18.	Incidental Services	16.1	• Not Applicable
19.	Spare Parts	17.1	Not applicable.

20.	Warranty of Goods	18.1	1 months from date of acceptance of the Goods.
21.	Period for the Replacement of Defective Goods	18.4	**7** days.
22.	Payment	22.1, 22.3& 22.4	The method and conditions of payment to be made to the Supplier under this Contract shall be as follows: Payment for Goods supplied from abroad: Payment of foreign currency portion shall be made in: (i) Advance Payment (ADVANCE_PAYMENT_PERCENT_HEALTH) percent of the Contract Price shall be paid within thirty (30) days of signing of the Contract, and upon submission of claim and a bank guarantee for equivalent amount valid until the Goods are delivered, and in the form, provided in the Tendering Documents or another form acceptable to the Purchaser. (ii) On Shipment: 0 percent of the Contract Price of the Goods shipped shall be paid through irrevocable confirmed letter of credit opened in favor of the Supplier in a bank in its country, upon submission of documents specified in GCC 13.1. (iii) On Acceptance: 100 percent of the Contract Price of Goods received shall be paid within thirty (30) days of receipt of the Goods upon submission of claim supported by the acceptance certificate issued by the Purchaser.

			Payment of local currency portion shall be made in **0** within thirty (30) days of presentation of claim supported by a certificate from the Purchaser declaring that the Goods have been delivered and that all other contracted Services have been performed. Payment for Goods and Services supplied from within the United Republic of Tanzania: Payment for Goods and Services supplied from within the United Republic of Tanzania shall be made in Tanzanian Shillings, as follows: (i) Not Applicable Percent of the Contract Amount. (ii) On Delivery: 0 percent of the Contract Price shall be paid on receipt of the Goods and upon submission of the documents specified in GCC 13.3. On Acceptance: 100 percent of the Contract Price shall be paid to the Supplier within thirty (30) days after the date of the acceptance certificate for the respective delivery issued by the Purchaser.
23	Currency of Payment	22.4	The Tanzanian Shilling
24.	Prices Adjustments	23.2	**NOT_APPLICABLE**
25.	Liquidated Damages and Maximum deduction of liquidated damages	29.1	In the case where the Performance Securing Declaration has been used, the rate of liquidated damages shall be 0.1 percent of the cost of undelivered goods for every day of delay to a maximum of

			10% of the contract sum.
26.	Settlement of Disputes	34.3	**Tanzania Institute of Arbitrators (TIArb)** **MBEYA**.
27.	Notices to PE	35.1	**MBEYA REGIONAL SECRETARIAT** **P.O BOX 754, MBEYA**

Special Conditions of Contract

PHARMACEUTICALS

(Additional Clauses)

The below data should be included in the **SCC** used in Tendering Documents for the procurement of pharmaceuticals.

SCC Number	**Information/Data Required**	**GCC Number**	**Amendments of, and Supplements to, Clauses in the GCC**
1.	Delivery and Documents	13.1 & 13.3	For Goods supplied from abroad: (ii) One original of the Certificate of Pharmaceutical Product as recommended by the WHO for each of the items supplied. (iii) Certificate of quality control test results in conformity with the World Health Organization "Certification Scheme on the Quality of Pharmaceutical Products Moving in International Trade" stating quantitative assays, chemical analysis, sterility, pyrogen content, uniformity, microbial limit, and other tests as appropriate to the Goods. (iv) Original copy of the certificate of weight issued by the port authority / licensed authority and six copies.

Special Conditions of Contract
VACCINES

(Additional Clauses)

The below data should be included in the **SCC** used in Tendering Documents for the procurement of vaccines.

SCC Number	**Information/Data Required**	**GCC Number**	**Amendments of, and Supplements to, Clauses in the GCC**
1.	Delivery and Documents	13.1 & 13.3	***For Goods supplied from abroad:*** (ii) One copy of the Lot Release Certificate issued by the NCA of the country of manufacture for each lot shipped. (iii) Certificate of quality control test results in conformity with the World Health Organization "Certification Scheme on the Quality of Pharmaceutical Products Moving in International Trade" stating quantitative assays, chemical analysis, sterility, pyrogen content, uniformity, microbial limit, and other tests as appropriate to the Goods. (iv) Original copy of the certificate of weight issued by the port authority / licensed authority and six copies.
			For Goods from within the United Republic of Tanzania:
			One copy of the Lot Release Certificate issued by the NCA of the country of manufacture for each lot shipped.
2.	Warranty of Goods	18.1	The Purchaser reserves the right to request evidence of bio-availability and/ or bio-equivalence data and / or evidence of the basis

			for expiration dating and other stability data concerning the goods to verify shelf life claimed for the Goods. If an adverse event following immunization (AEFI) occurs in the United Republic of Tanzania and the cause of such event cannot be immediately established, the Purchaser will, with all urgency and in accordance with the procedures laid down by the NCA of the United Republic of Tanzania, take steps to advise the supplier in order that an investigation may be launched immediately. If the vaccine has been supplied through an agency of the United Nations, the most current procedures laid down by the WHO for such situations will be used.

Special Conditions of Contract
CONDOMS

The following **SCC** shall supplement the **GCC** in the procurement of condoms. Whenever here is a conflict, the provisions herein shall prevail over those in the **GCC**. The corresponding clause number of the **GCC** is indicated in parent theses.

S/N	Information/Data Required	GCC Number	Amendments of, and Supplements to, Clauses in the GCC
1.	Inspection and Tests	11.1 & 11.2	(a) The Supplier shall test batches of Goods ready for shipment in accordance with the WHO specification. The size of the sample for testing will be calculated by reference to ISO 2859-1. With each consignment, the Supplier must provide a certificate of quality control test results in conformity with the standards laid down in ISO 2859-1 and in accordance with the general sampling levels appropriate to each feature as necessary. The Supplier will bear the cost of such tests.
2.	Delivery and Documents	13.1 & 13.3	For Goods supplied from abroad: (ix) original copy of quality control tests for each consignment as stated in SCC 9 above. (x) original copy of the certificate of inspection furnished to Supplier by nominated inspection agency and six copies. [where separate inspection is required] For Goods from within the United Republic of Tanzania: i) Certificate of in-house analysis.

Section X: Contract Forms

This Section contains forms that, once completed, will form part of the Contract. The forms for Performance Security/Performance Securing Declaration and Advance Payment Security, when required, shall only be completed by the successful Tenderer after contract award. The section also contains the Letter of Intention to Award the Contract, which shall not form part of the contract.

Notice of Intention to Award a Contract

[*Letterhead paper of the PE*]

Ref No: *[insert ref. no.]*..... Date: .....

To: . . . . . ***[name and address of the Supplier]*** . . . . . .

RE: NOTIFICATION OF THE INTENTION TO AWARD CONTRACT NUMBER [*insert number of contracts***] FOR [***insert description***]**

Reference is made to the above subject matter. The submitted tenders were evaluated according to the criteria stated in the tender document. In accordance with the requirements of Public Procurement Act, Cap 410 , we announce our intention to award a contract to M/s*: (Insert the name of the firm)* for a contract price of *(insert the contract award price and currency)* and for a completion period/delivery period of *(insert the duration)*

Your tender was not considered for the award of the contract due to the following reasons:-

1) .....

2) .....

3) .....

Be informed that, you have five (5) working days from the date of this letter, within which to submit any complaints you may have regarding this award decision and/or circumstances surrounding the rejection of your tender for administrative review,. The complaints must be in writing, clearly identifying the tender in question, detailing ground(s) of the complaint and should be submitted to *(insert the title of Accounting Officer)* through NeST.

We appreciate your interest in doing business with us and encourage you to participate in our future tenders.

Authorized Signature: .....

Name and Title of Signatory: .....

Name of PE: .....

1 *Insert the reasons for non-selection of the tenderer for the award of contract. The reasons given here should be those which appears in the evaluation report and which were approved by the Tender Board as justifiable reasons to turn down the offer given by the tenderer.*

Letter of Acceptance

[Letterhead paper of the PE]

[date]

To: *[name and address of the Supplier]*

RE: NOTIFICATION OF AWARD OF CONTRACT FOR TENDER NO.[insert tender number] **FOR** [insert tender description]

This is to notify you that your tender dated *[insert date]* for execution of the Contract Number *[insert Contract number and description, as given in the Special Conditions of Contract]* for the Accepted Contract Amount of the equivalent of *[insert amount in numbers and words and name of currency]*, as corrected and modified in accordance with the Instructions to Tenderers is hereby accepted by us.

You are requested to furnish the Performance Securing Declaration or Performance Security within 14 days in accordance with the Conditions of Contract, using for that purpose the Performance Security Form included in Section X, Contract Forms, of the Tendering Document.

Authorized Signature: .....

Name and Title of Signatory: .....

Name of Agency: .....

Attachment: Contract Agreement

Copy:, PPRA, CAG, Office of the Attorney General, GAMD, IAG, TRA and Adjudicator's Appointing Authority,

Insert the appropriate form of security to be furnished. The Performance Securing Declaration shall only be applicable for Tenders falling under exclusive preference.

Form of Contract

THIS CONTRACT (hereinafter referred as the ‘‘Contract’’) is made this [*day of the month*] day of [*insert a month*], [*insert a year*]_ between *[name and address of Purchaser]* (hereinafter referred as “the Purchaser”) of the one part and *[name and address of Supplier]* (hereinafter referred as “the Supplier”) of the other part:

[***Note****: In the text below, text in brackets is optional; all notes should be deleted in final text. If the Supplier consist of more than one Entity, the above should be partially amended to read as follows:*]

“*[insert the name of Purchaser]* (hereinafter referred as the “Purchaser”) and, on the other hand, a joint venture/consortium/association consisting of the following entities namely, *[insert of name of entity]* and *[insert name of entity] and [etc.]* (hereinafter referred as the “Supplier”) each of which shall be jointly and severally liable to the Purchaser for all the Supplier's obligations under this Contract.

WHEREAS the Purchaser invited Tenders for certain goods and ancillary services, *viz.,* [*insert brief description of goods and services*] and has accepted a Tender by the Supplier for the supply of those goods and services in the sum of [*insert contract price in words and figures*](hereinafter called "the Contract Price").

NOW THIS CONTRACT WITNESSETH AS FOLLOWS:

1. In this Contract words and expressions shall have the same meanings as are respectively assigned to them in the Conditions of Contract referred to.
2. The following documents shall constitute the Contract between the Purchaser and the Supplier, and each shall be read and construed as an integral part of the Contract:

   (a) Form of Contract,

   (b) Letter of Acceptance

   (c) Form of Tender

   (d) Special Conditions of Contract,

   (e) General Conditions of Contract,

   (f) Specifications

   (g) Completed Schedules (including Price Schedules),

   (h) Performance Security;

   (i) Power of Attorney; and

(j) [Other relevant document(s): [*List any such as negotiation minutes if applicable*]

3. The mutual rights and obligations of the Purchaser and the Supplier shall be as set forth in the Contract, in particular:

   a. In consideration of the payments to be made b y the Purchaser to the Supplier as hereinafter mentioned, the Supplier hereby covenants with the Purchaser to provide the Goods and Services and to remedy defects therein in conformity in all respects with the provisions of the Contract.

   b. The Purchaser hereby covenants to pay the Supplier in consideration of the provision of the Goods and Services and the remedying of defects therein, the Contract Price or such other sum as may become payable under the provisions of the Contract at the times and in the manner prescribed by the Contract.

IN WITNESS WHEREOF, the Parties hereto have caused this Contract to be executed in their respective names as of the day, month and year specified above.

SIGNED FOR AND ON BEHALF OF:

THE PURCHASER	**THE SUPPLIER**
Name: ..... (*Authorized Representative*)	**Name:** ..... (*Authorized Representative*)
Designation:	**Designation:** .....
Signature:	**Signature:** .....
WITNESS	**WITNESS**
Name:	**Name:** .....
Designation:	
Signature:	

Designation:

.....

Signature:

.....

PERFORMANCE SECURING DECLARATION

Date: *[insert **date** (as day, month and year)]*

Contract N o .: *[insert **Contract number]***

To:*[insert **complete name of Purchaser**]*

I/We, the undersigned, declare that:

1. I/We understand that, according to your conditions, to guarantee the faithful performance by the Contractor of its obligations under the Contract, I/We shall submit this form of Performance Securing Declaration within a maximum period of fourteen (14) calendar days from the date of the Letter of Acceptance and prior to the signing of the Contract.
2. I/We accept that: I/We will be disqualified from tendering for any procurement contract with any procuring entity for t h e p e ri o d o f ti m e determined by the Public Procurement Regulatory Authority in accordance with the procedures stipulated in the Public Procurement Act and Public Procurement Regulations if I/We have failed to execute the Contract in accordance with the Term and Conditions therein.

I/We understand that this Performance Securing Declaration shall cease to be valid upon satisfactory performance and final acceptance of the goods by the Purchaser.

Signed: *[insert signature of person whose name and capacity are shown]* in the capacity of *[insert legal capacity of person signing the Performance Securing Declaration]*

Name: *[insert **complete name of person signing the Performance Securing Declaration**]*

Duly authorised to sign the Contract for and on behalf of: *[insert **complete name of Supplier]***

Dated on day of _, *[insert **date of signing**]*

Corporate Seal (where appropriate)

Used as an alternative performance security for Contracts whose value falls under the Regional Exclusive Preference. It shall be submitted within fourteen (14) days after receiving the Letter of Acceptance

Performance Security Form

PERFORMANCE BANK GUARANTEE
(UNCONDITIONAL)

[*Bank/successful tenderer providing the guarantee shall complete this form according to the instructions given in the brackets, if Procuring Entity requires this type of security*.]

[*Insert name of bank, and address of the issuing branch or office*]

Beneficiary: [*Insert name and address of the Procuring Entity*]

Date: [*Insert date*]

PERFORMANCE GUARANTEENUMBER: [*Insert Performance Guarantee number*]

We have been informed that[*insert name of the Supplier/Service Provider*] (hereinafter referred to as "*Supplier/Service Provider*") has been awarded the contract under Tender No. *[insert the tender reference number]* from your institution for the execution of *[insert the name and brief description of the tender*]."

Furthermore, we understand that, according to the terms of the award of the contract, a performance guarantee is required.
At the request of the Supplier/Service Provider, we [*insert name of the Bank*]hereby undertake to pay you any sum or sums not exceeding in total an amount of[*insert amount in figures and in words*], payable in the type and proportion of currency in which the contract price is payable, upon receipt of your written demand accompanied by a written statement indicating the Supplier's/Service Provider's failure to fulfil its obligations under the contract and a copy of the contract signed between the Supplier/Service Provider and your institution, without the need to prove or to show grounds for your demand or the amount specified therein.

This guarantee shall expire no later than 28 days from the date of issuance of the certificate of completion of a contract, calculated based on a copy of the certificate that we shall be provided with, or on *[insert day]* of *[insert month], [insert year],* whichever occurs first. Consequently, any demand for payment under this guarantee must be received by us at this office on or before that date.

…….

[Signature of the Bank's authorised representative(s)]

Note: All italicized text (including footnotes) is for use in preparing this form and shall be deleted from the final product.

Performance Bond

[Guarantor letterhead]

By this Bond *[insert name of Principal]* as Principal (hereinafter called "the Supplier") and *[insert name of Surety]* as Surety (hereinafter called "the Surety"), are held and firmly bound unto *[insert name of Purchaser]* as Obligee (hereinafter called "the Purchaser") in the amount of *[insert amount in words and figures]*, for the payment of which sum well and truly to be made in the types and proportions of currencies in which the Contract Price is payable, the Supplier and the Surety bind themselves, their heirs, executors, administrators, successors and assigns, jointly and severally, firmly by these presents.

WHEREAS the Supplier has entered into a written Agreement with the Purchaser dated the day of , 20 , for *[name of contract and brief description of Works]* in accordance with the documents, plans, specifications, and amendments thereto, which to the extent herein provided for, are by reference made part hereof and are hereinafter referred to as the Contract.

NOW, THEREFORE, the Condition of this Obligation is such that, if the Supplier shall promptly and faithfully perform the said Contract (including any amendments thereto), then this obligation shall be null and void; otherwise, it shall remain in full force and effect. Whenever the Supplier shall be, and declared by the Purchaser to be, in default under the Contract, the Purchaser having performed the Purchaser's obligations thereunder, the Surety may promptly remedy the default, or shall promptly:

(1) complete the Contract in accordance with its terms and conditions; or

(2) obtain a Tender or tenders from qualified Tenderers for submission to the Purchaser for completing the Contract in accordance with its terms and conditions, and upon determination by the Purchaser and the Surety of the lowest responsive Tenderer, arrange for a Contract between such Tenderer and Purchaser and make available as work progresses (even though there should be a default or a succession of defaults under the Contract or Contracts of completion arranged under this paragraph) sufficient funds to pay the cost of completion less the Balance of the Contract Price; but not exceeding, including other costs and damages for which the Surety may be liable hereunder, the amount set forth in the first paragraph hereof. The term "Balance of the Contract Price," as used in this paragraph, shall mean the total amount payable by Purchaser to Supplier under the Contract, less the amount properly paid by Purchaser to Supplier; or

(3) pay the Purchaser the amount required by the Purchaser to complete the Contract in accordance with its terms and conditions up to a total not exceeding the amount of

this Bond.

The Surety shall not be liable for a greater sum than the specified penalty of this Bond.

Any suit under this Bond must be instituted before the expiration of one year from the date of issue of the Certificate of Completion.

No right of action shall accrue on this Bond to or for the use of any person or corporation other than the Purchaser named herein or the heirs, executors, administrators, successors, and assigns of the Purchaser.

In testimony whereof, the Supplier has hereunto set his hand and affixed his seal, and the Surety has caused these presents to be sealed with his corporate seal, duly attested by the signature of his legal representative, this day of 20.

SIGNED ON behalf of

By in the capacity of

In the presence of

SIGNED ON behalf of

In the capacity of

In the presence of

Bank Guarantee Form for Advance Payment

Date: [*insert date*]

IFT: [*insert name and number of IFT*]

Contract*:* [*insert name and number of Contract]*

*To: [**insert name and address of Purchaser**]*

Dear Sir or Madam

In accordance with the payment provision included in the Special Conditions of Contract (SCC), which amends GCC 23 to provide for advance payment, *[name and address of Supplier]* (hereinafter called "the Supplier") shall deposit with the Purchaser a Bank Guarantee to guarantee its proper and faithful performance under the said Clause of the Contract in an amount of *[amount of guarantee in figures and words]*.

We, the *[bank or financial institution]*, as instructed by the Supplier, agree unconditionally and irrevocably to guarantee as primary obligator and not as surety merely, the payment to the Purchaser on its first demand without whatsoever right of objection on our part and without its first claim to the Supplier, in the amount not exceeding *[amount of guarantee in figures and words]*.

We further agree that no change or addition to or other modification of the terms of the Contract to be performed thereunder or of any of the Contract documents which may be made between the Purchaser and the Supplier, shall in any way release us from any liability under this guarantee, and we hereby waive notice of any such change, addition, or modification.

This guarantee shall remain valid and in full effect from the date of the advance payment received by the Supplier under the Contract until *[date]*.

Yours truly,

Signature and seal of the Guarantors

_____

[name of bank or financial institution]

_____

[address]

_____

[date]

Sample form for Advance Payment Application

Insert[Tenderer Logo]

Insert [TENDERER_NAME]

Insert [TENDERER_ADDRESS]

APPLICATION FOR ADVANCE PAYMENT

A. Contract Data

This application is made under General Conditions of Contract (GCC) Clause [*insert relevant clause*]

S/N	DESCRIPTION	PARTICULARS
1.	Name of Procuring Entity:	*Insert [PE_NAME]*
2.	Contractor's/Vendor's Name:	*Insert[TENDERER_NAME]*
3.	Contract No:	*Insert [CONTRACT_NUMBER]*
4.	Contract Description:	*Insert[TENDER_DESCRIPTION]*
5.	Contract Signing Date:	*Insert [CONTRACT_SIGNING_DATE]*
6.	Start Date:	*Insert [CONTRACT_START_DATE]*
7.	Intended Completion Date:	*Insert [INTENDED_COMPLETION_DATE]*
8.	Application for Payment No:	*Insert [APPLICATION_NUMBER]* *First Application* *Second Application* *Third Application etc.*
9.	Application Date:	*Insert*

		[ADVANCE_PAYMENT _APPLICATION_DATE]
10.	Contract Price:	*Insert [CONTRACT_PRICE]*
11.	Advance Payment Percent:	*Insert [ADVANCE_PAYMENT_PERCENT]*
12.	Advance Payment Currency:	*Insert [ADVANCE_PAYMENT_CURRENCY]* *(Currency One)* *(Currency Two)* *(Currency Three)*
13.	Advance Payment Amount:	*Insert [ADVANCE_PAYMENT_AMOUNT]* *Payment for Currency One* *Payment for Currency Two* *Payment for Currency Three*
14.	Tenderer Bank Details:	Tenderer Account Number: Tenderer Account Name: Bank Name: Bank Region: Bank Country: Bank SWIFT Code:

B. Commitment by *Contractor, Consultant, Supplier or Service Provider*

The advance payment shall only be used for the purpose of mobilization expenses or other requirements specifically for execution of the assignment as stipulated in the contract document and we shall demonstrate to the Project Manager that the advance payment has been used for the intended purposes.

Applied by:(Insert name of representative of the Contractor, Consultant, and Supplier or Service provider)

Signed by:*(insert name)*

Signature:*(insert signature)*

Title:*(insert title)*