

MAHARASHTRA MEDICAL GOODS
PROCUREMENT AUTHORITY

Tender for Supply of Essential Medicines
(Year 2025-26)

Tender reference No: E-248/MMGPA/Essential Medicines (2025-26)

1st Floor, Aarogya Bhawan
St. George's Hospital Compound,
Near C.S.M.T. Railway Station, Mumbai - 400 001.
Maharashtra

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Table of Contents

Table of Contents	2
Disclaimer	5
Abbreviations and Glossary	7
Bid Schedule	10
Fact Sheet.....	12
1. Section 1: Introduction.....	13
2. Section 2: General Definitions	15
3. Section 3: General Terms and Conditions.....	18
3.1. Responsibility of verification of contents of Tender Document	18
3.2. Authorized Signatory for the Tender Document.....	18
3.3. Period of Validity of Bid	18
3.4. Earnest Money Deposit (EMD)	18
3.5. Submission of Bids	19
3.6. Language	23
3.7. Format of Bid.....	23
3.8. Number of Bids and Cost of bidding:.....	23
3.9. Amendment of Tender document	23
3.10. Acknowledgement by Bidder	24
3.11. Right to accept or reject any or all Bids.....	24
3.12. Pre-Bid Meeting.....	25
3.13. Clarifications post Pre bid Meeting	26
3.14. Modification/substitution/withdrawal of Bids	26
3.15. Proprietary data	26
3.16. Correspondence with the Bidder	26
3.17. Opening and Evaluation of Bids	27
3.18. Confidentiality	28
3.19. Clarifications regarding Evaluation	28
3.20. Selection of Bidder	29
3.21. Unethical Practices	30
3.22. Code of Integrity	31
3.23. Payment Provisions	32
3.24. Termination of Contract.....	32
3.25. Dispute Resolution.....	34
3.26. Governing law and jurisdiction.....	34
3.27. Indemnification.....	34
3.28. Saving clause.....	34
3.29. Force Majeure	35

4. Section 4: Specific Terms and Conditions	35
4.1. Eligibility Criteria and Supporting Documents to be Submitted	35
4.2. Manufacturing/Importing and Product Permissions.....	43
4.3. Minimum Tender Quantity	43
4.4. Market Standing.....	44
4.5. Inspection of Manufacturing Facilities	45
4.6. Shelf Life of the Drugs	46
4.7. Method of Placing Purchase Orders.....	46
4.8. Execution of Contract	46
4.9. Performance Security	46
4.10. Supply Conditions	47
4.11. Packing and Labeling	48
4.12. Schedule for packaging of essential drugs	49
4.13. Aggregation and “parent-child” relationship serialisation.....	52
4.14. Deduction & Damages on Quality Failure.....	53
4.15. Blacklisting of Product/Tenderer on withdrawal of Tender	55
4.16. Blacklisting for Non-Supply of purchase Order(s)	55
4.17. Blacklisting of Product(s) based on Quality Test by the Empanelled Laboratories.....	55
4.18. Blacklisting of Product(s) based on Quality Test by Statutory Authorities.....	56
4.19. Blacklisting of the Supplier on Quality Failure.....	57
4.20. Procedure for Blacklisting/Debarment:.....	57
Appendix-A: Schedule of Requirements	58
Appendix-B: Checklist	118
Annexure 1: Technical Specifications and Compliance.....	120
Annexure-2: Letter Comprising Technical Bid.....	122
Annexure-3: Proforma for Production and Sale Statement.....	125
Annexure-4: Details of Manufacturing Unit	126
Annexure-5: Details of Items Quoted with Drug Code.....	130
Annexure 6: Annual Turnover Statement for Three Years	131
Annexure-7: Format for Power of Attorney for signing of Bid	132
Annexure-8: Affidavit for blacklisting	134
Annexure- 9: Litigation Affidavit	135
Annexure-10: Mandate Form	136
Annexure-11: Indicative format for PRICE BID (BOQ) to be submitted online only	138
Annexure-12: Statement showing comparative prices offered by the tenderer in other tenders of the same product	139
Schedule 1: Contract Form.....	140
Schedule-2 Performance Security Form.....	143

Schedule-3: Supply Schedule.....	145
Schedule 4: Schedule for Packing of Drugs and Medicines	146
Schedule-5: Bar Code & Advance Shipment Notification details	Error! Bookmark not defined.

MMGPA Tender

Disclaimer

The information contained in this Tender Document or subsequently provided to Bidder(s), whether verbally or in documentary or any other form, by or on behalf of the Maharashtra Medical Goods Procurement Authority or any of its employees or advisors, is provided to Bidder(s) on the terms and conditions set out in this Tender Document subject to which such information is provided.

This Tender Document is not an agreement and is neither an offer nor invitation by the MMGPA to the prospective Bidders or any other person. The purpose of this Tender Document is to provide interested parties with information that may be useful to them in making their financial offers (Bids) pursuant to this Tender Document. This Tender Document includes statements, which reflect various assumptions and assessments arrived at by the MMGPA in relation to the project. Such assumptions, assessments and statements do not purport to contain all the information that each Bidder may require. This Tender Document may not be appropriate for all persons, and it is not possible for the MMGPA, its employees or advisors to consider the investment objectives, financial situation and particular needs of each party who reads or uses this Tender Document. The assumptions, assessments, statements and information contained in this Tender Document may not be complete, accurate, adequate or correct. Each Bidder should, therefore, conduct its own investigations and analysis and should check the accuracy, adequacy, correctness, reliability and completeness of the assumptions, assessments, statements and information contained in this Tender Document and obtain independent advice from appropriate sources.

Information provided in this Tender Document to the Bidder(s) is on a wide range of matters, some of which may depend upon interpretation of law. The information given is not intended to be an exhaustive account of statutory requirements and should not be regarded as a complete or authoritative statement of law. The MMGPA accepts no responsibility for the accuracy or otherwise for any interpretation or opinion on law expressed herein.

The MMGPA, its employees and advisors make no representation or warranty and shall have no liability to any person, including any Bidder under any law, statute, rules or regulations or tort, principles of restitution or unjust enrichment or otherwise for any loss, damages, cost or expense which may arise from or be incurred or suffered on account of anything contained in this Tender Document or otherwise, including the accuracy, adequacy, correctness, completeness or reliability of the Tender Document and any assessment, assumption, statement or information contained therein or deemed to form part of this Tender Document or arising in any way for participation in this Tender Document.

The MMGPA also accepts no liability of any nature whether resulting from negligence or otherwise howsoever caused arising from reliance of any Bidder upon the statements contained in this Tender Document.

The MMGPA may, in its absolute discretion but without being under any obligation to do so, update, amend or supplement the information, assessment or assumptions contained in this Tender Document.

The Bidder shall bear all its costs associated with or relating to the preparation and submission of its Bid including but not limited to preparation, copying, postage, delivery fees, expenses

associated with any demonstrations or presentations which may be required by the MMGPA or any other costs incurred in connection with or relating to its Bid. All such costs and expenses will remain with the Bidder and the MMGPA shall not be liable in any manner whatsoever for the same or for any other costs or other expenses incurred by a Bidder in preparation or submission of the Bid, regardless of the conduct or outcome of the bidding process.

MMGPA Tender

Abbreviations and Glossary

Abbreviations and Acronyms	Description
BET	Bacterial Endotoxin Test
BG	Bank Guarantee
BIS	Bureau of Indian Standards
BOQ	Bill of Quantity/(ies)
BP	British Pharmacopeia
CA	Chartered Accountant
CLA	Central Level Authority
COPP	Certificate of Pharmaceutical Products
CT	Clinical Trials
DCGI	Drug Controller General of India
DMER	Directorate of Medical Education & Research
DPCO	Drug Price Control Order
DPIIT	Department for Promotion of Industry and Internal Trade
EMD	Earnest Money Deposit
ESIC	Employees State Insurance Corporation
FDA	Food and Drug Administration
FEMA	Foreign Exchange Management Act 1999
GMP	Good Manufacturing Practices
G.R.	Government Resolution
GST	Goods and Services Tax
GS1	Global Standards1
HBPCL	Haffkine Bio-Pharmaceutical Corporation Limited
IEC	Import Export Code
IP	Indian Pharmacopeia
ISI	Indian Standards Institute
I.V.	Intra Venous
LLP	Limited Liability Partnership
LCBS	Least Cost Based Selection
MCGM	Municipal Corporation of Greater Mumbai
MDR	Medical Device Rules
MMGPA	Maharashtra Medical Goods Procurement Authority
MRP	Maximum Retail Price
MSC	Market Standing Certificate
MSEs	Micro and Small Enterprises
NABL	National Accreditation Board for Laboratories
NEFT	National Electronic Funds Transfer
NSQ	Not of Standard Quality
NPPA	National Pharmaceutical Pricing Authority
OEM	Original Equipment Manufacturer
PAN	Permanent Account Number
PoA	Power of Attorney

Abbreviations and Acronyms	Description
PVC	Poly Vinyl Chloride
PVdC	Poly Vinylidene Chloride
RFP	Request For Proposal
RTGS	Real Time Gross Settlement
SLA	State Level Authority
SSI	Small Scale Industries
TAA	Tender Accepting Authority
TIA	Tender Inviting Authority
UDIN	Unique Document Identification Number
USP	United States Pharmacopeia
WHO	World Health Organization

MAHARASHTRA MEDICAL GOODS PROCUREMENT AUTHORITY

Bid Notice

**Tender reference No: E-248/MMGPA/Essential Medicines
(2025-26)**

Chief Executive Officer, Maharashtra Medical Goods Procurement Authority, Mumbai invites **ONLINE BID** for the year **2025-26** in **two envelope system** from the manufacturers/importers for the purchase of following items.

S. No.	Description	Tender Fee (Rs.)	EMD (Rs.)
1.	As per Appendix– A	47,000/-	Subject to Clause 3.4, a Bidder is required to furnish/pay a minimum EMD amount of Rs.2.00 lacs. However, if the Bidder is submitting its quote for more than 8 drugs, then for each additional drug beyond 8 number of drugs, the EMD amount shall correspondingly increase by a value of Rs. 25,000/- per additional drug, subject to a maximum EMD amount of Rs. 3.00 lacs.

Bid Schedule

All bid related activities (process) like downloading of bid document, submission of bid and submission of EMD and other documents will be governed as per the time schedule given below:

Sr. No.	Activity	Period
1	Period of sale of Tender Document/ download	From – 08-12-2025, 05:30 PM
2	Date for submission of queries	Before pre-bid meeting
3	Date of pre-bid meeting	15.12.2025 at 2:00 PM To 03:00 PM (Bidder should have to submit queries in the prescribed format, through email or hard copy before scheduled time of meeting to be held at Ground Floor, Aarogya Bhawan, Commissionerate of Health Services, Mumbai 400001)
4	E- tender submission duration	From- 08.12.2025 05:30 PM to 19.12.2025 up to 12:00 PM
5	Last date and time for submission of Bid: (Bid Due Date)	19.12.2025 up to 12:00 PM
6	Date and time of opening of Technical Bid (envelope No.1)	22.12.2025 at 02:00 PM
7	Date and time of opening of Price Bid (envelope No. 2)	To be notified to the technically qualified Bidders.
8	Validity of Tender	180 days from the Bid Due Date

Address for communication : 1st Floor, Aarogya Bhawan,
St. Georges Hospital Compound,
Near CSMT Railway Station, Mumbai- 400 001.

Tender Document is available to be downloaded from the Official Website (<https://mahatenders.gov.in>), however at the time of submission of Bid, any interested eligible bidder would be required to pay by online payment of a non-refundable tender fee of INR 50,000 and enclose a copy of the receipt of payment with the technical bid. The payment of the Tender fee shall be made only online through payment gateway in **A/c of "Maharashtra Medical Goods Procurement Authority, Mumbai"**.

As per Revised Manual of Office Procedures for Procurement by the Government Department issued through Government Resolution no. Bha.kha.sa-2014/ Pra. Kra. 82/Section-III/Industry-4 by Department of Industry , Energy & Labour, Government of Maharashtra, dated 01.12.2016 – Bidder including Government Boards/Corporation/Undertakings and manufactures/ suppliers who are MSEs

registered under “Micro, Small and Medium Enterprises Development Act 2006” are exempted from paying Tender Fee and Earnest Money Deposit.

Non-payment of Earnest Money Deposit, unless exempted herein, will result in the rejection of the bid summarily without any notice. The conditional bid shall be rejected.

Chief Executive Officer, Maharashtra Medical Goods Procurement Authority, Mumbai reserves all the rights regarding this Tender Document and the procedure outlined therein

**Sd/-
CHIEF EXECUTIVE OFFICER,
MAHARASHTRA MEDICAL
GOODS PROCUREMENT
AUTHORITY MUMBAI**

Fact Sheet

Clause Reference	Topic
Price Bid Evaluation	<i>The method of selection is LCBS (Least Cost Based Selection-L1).</i>
Downloading Tender Document	Tender Document can be downloaded from https://mahatenders.gov.in and from website of MMGPA for reference (“ Official Websites ”).
Earnest Money Deposit (EMD)	Bidders are required to pay the EMD/Bid Security as per the Clause 3.4. through online mode on https://mahatenders.gov.in .
Scope of Work	Supply of essential medicines (2025-26) in accordance with terms of this Tender Document.
Pre-bid meeting and clarifications	A pre-bid meeting will be held on dt. 15.12.2025 02:00 PM , Meeting will be held in two sessions: A) Issues Regarding specifications, B) Issues Regarding overall tender documents The participants should have an authority letter of the company which they are representing. Maximum two representative will be allowed per firm / company. Only authorised participants inside the meeting hall. Bidder may submit queries through email or hard copy before scheduled time of pre-bid meeting.
Taxes	For all goods/services supplied, the Bidder shall be entirely responsible for bearing of all taxes, stamp duties, license fees, and other such levies imposed/incurred until delivery of the contracted products or services.
Bid Validity	Bids must remain valid till 180 days from the Bid Due Date.
Submission of Responses	Bidders must upload and submit all the documents on the e-tendering portal https://mahatenders.gov.in (“ E-Tender Portal ”). <i>Each of the documents must be uploaded in the format specified in this Tender Document.</i>
Submission of Bids	This is online process; interested Bidders are required to submit the Bids online by the date and time specified in the Tender Document.
Last Date of Submission	Bids submitted after Dt. 19.12.2025 12:00PM will not be accepted by the e-Tender portal.
Tender Fee	All Bidders shall pay Tender Fee of Rs.47,000

Section 1: Introduction

Maharashtra Medical Goods Procurement Authority (MMGPA), Mumbai has been formed as per the Maharashtra Medical Goods Procurement Authority Act 2023 (Mah. Act No. XIII of 2023) (“**Act**”). MMGPA works as a single point procurement of certain medical goods and execution of turnkey projects with highest standards of transparency, fairness, equity and for timely supply at optimum and uniform rates and of desired quality and quantity for the Public Health Department, Medical Education & Drug Department of Government of Maharashtra and other departments as mentioned in the said Act. To meet the objective of the Act, MMGPA in exercise of its powers under section 36 of the Act issued the regulations named, Maharashtra Medical Goods Procurement Authority Regulations, 2025 (“**Regulations**”) as amended from time to time.

- 1.1 **Chief Executive Officer, Maharashtra Medical Goods Procurement Authority, Mumbai**, hereinafter referred to as the (“**Purchaser**”) invites online Bid in two envelope systems for supply of item specified in ‘**Appendix-A** Schedule of Requirements’, for use in public health facilities in the State of Maharashtra. Any applicable orders/ circulars issued by Govt. of Maharashtra from time to time will be applicable to this bidding process.
- 1.2 All bid related activities (process) like Tender Document downloading, bid submission and submission of EMD and other documents will be governed by the bid schedule given in bid notice.
- 1.3 All activities of this bid are carried out **online** on website <https://mahatenders.gov.in>. The Tender Document is uploaded on Government of Maharashtra, (GoM) e-tendering website <https://mahatenders.gov.in> and has to be downloaded as well as filled up and submitted **online only**. The Bidders are required to submit online Tender Fees (non-refundable) as mentioned, through **online payment gateway in A/c of "Maharashtra Medical Goods Procurement Authority, Mumbai"**. In no case, the Tender Fee should be mixed with EMD amount. The Bid shall be liable to be rejected summarily upon failure to follow procedure prescribed in the Tender Document.
- 1.4 **The quantities mentioned in the Bid are only approximate estimated quantities (“Tendered Quantity”). The Chief Executive Officer, Maharashtra Medical Goods Procurement Authority, Mumbai reserves the right to increase or decrease the quantities’, maximum up to 50% of the quantities to be purchased without assigning any reason thereof.**
- 1.5 If any Bidder wishes to lodge any complaint regarding the evaluation of his Bid, the Bidder shall submit the complaint within 48 (forty-eight) hours from the declaration of Bidders whose Bids are technically qualified along with deposit of **Rs. 50,000 (Rupees Fifty Thousand only)** online in favour of **Chief Executive Officer, Maharashtra Medical Goods Procurement Authority, Mumbai** in the form of deposit. This complaint will be submitted to appropriate forum along with facts. The amount so deposited shall be refunded, if after scrutiny the complaint is found to be true by the appropriate forum. However, if the complaint is found to be false and malafide the deposit will be forfeited. No interest shall be paid against this deposit.

1.6 E-bidding process related Queries can be sent on email – eproc.support@maharashtra.gov.in /Help: The 24 x 7 Toll Free Telephonic Help Desk Number 1800-3070-2232. / Mobile: +91-7878107985 , +91-7878107986 , + 91-7878007972 and +91-7878007973

Sd/-
Chief Executive Officer,
Maharashtra Medical Goods
Procurement Authority Mumbai

MMGPA Tender

Section 2: General Definitions

- 2.1. Applicable Laws:** shall mean all laws and regulations brought into force and effect by GOI or the State Government of Maharashtra or Food and Drug Administration including the CDSCO norms, Drugs and Cosmetics Act, 1940, Medical Device Rules, 2017 (MDR), NPPA, CVC Guidelines, New Drugs and Clinical Trials Rules, 2019, Drug Price Control Orders, rules, regulations, notifications, directives, policies and office memorandums, made thereunder and as amended from time to time and judgements, decrees, injunctions, writs and orders of Hon'ble Supreme Court or High Court, applicable to this Tender Document and the exercise, performance and discharge of the respective rights and obligations of the parties hereunder, as may be in force and effect during the tenure of the Tender Document or Contract.
- 2.2. Bid / Tender:** shall mean the two-envelope submitted i.e., envelope no. 1 (Technical Bid) and envelope No. 2 (Price Bid), collectively.
- 2.3. Blacklisting/Debaring:** The occurrence of any event, as mentioned hereinbelow, shall make the Bidder ineligible from participating in the future bids of Purchaser for a period of 2 to 5 years, as specified in the blacklisting order. The period of Blacklisting/Debaring shall be decided on the basis of number/nature of violations of the tender conditions and the loss/hardship caused/likely to be caused to the Purchaser on account of each such violations, generally relating to:
- a) supply of substandard, misbranded, adulterated or spurious or any drugs/products manufactured/imported in contravention of any of the laws of the land and/ or,
 - b) indulging in unethical practices as detailed in Clause 3.21 and/ or,
 - c) indulging in fraudulent practices at the time of making the bid or at any time during the validity of the tender or the contract thereof. The event will include, among all other things, making false/misleading declarations statements, presenting false/ misleading/ fabricated/ forged document(s), trying to influence/ affect/ stall the tender/ procurement/ payment processes in anyway, making false/ baseless complaints about other bidders or bids or drugs or any person/organization/related to the tender activities etc., and such activities as specified in the Tender Document.
 - d) Any supplied drug/medicines if declared as Not of Standard Quality (NSQ) by the competent authority during the statutory sampling of the relevant drug/medicines shall also be ground for blacklisting by the Purchaser.
- 2.4. BOQ:** Bill of Quantity or the Schedule of Quantity in which the rates are to be filled in by the Bidder or commonly called the price bid.
- 2.5. Contract:** A Contract for the supply of an approximate quantity of item(s) at a specified price and period as mentioned in Purchase Order(s) issued by the Purchaser from time to time during the Period of Contract.
- 2.6. Drug Inspector:** As defined under the Drugs and Cosmetics Act, 1940, including any amendments thereof.
- 2.7. Empaneled Laboratory:** Drug Testing Laboratory approved under the Drugs and Cosmetics Act 1940 and accredited by NABL, selected by the Purchaser either

through open tender process or by expression of interest or otherwise for the purpose of conducting analytical testing/evaluation of drugs procured through the tender.

- 2.8. Government:** means the Central Government or the Government of Maharashtra, as the case may be, and includes agencies and public sector enterprises under it, in specific contexts.
- 2.9. MDR, 2017 – Medical Device Rules** published under sub-section (1) of Section 12 and Sub-section (1) of Section 33 of the Drugs and Cosmetics Act, 1940 (23 of 1940), in the Gazette of India to regulate the import, manufacture, distribution and sale of Medical Devices and any subsequent amendments thereto.
- 2.10. Market Standing Certificate:** A Market Standing Certificate (MSC) is a document that verifies a company's good standing and track record in the pharmaceutical or medical device industry issued by State Drug Control Authority or Central Drugs Standard Control Organization (CDSCO).
- 2.11. New Drugs and Clinical Trials Rules, 2019-** Rules published under sub-section (1) of Section 12 and Sub-section (1) of Section 33 of the Drugs and Cosmetics Act, 1940 (23 of 1940) and any subsequent amendments thereto, in the Gazette of India which shall apply to all new drugs, investigational new drugs for human use, clinical trial, bioequivalence study, bioavailability study and ethics committee.
- 2.12. Notification of Award (NoA):** is an intimation informing the successful Bidder, the approximate quantity for which the Tender Document is awarded and requiring the Bidder to execute an agreement in the prescribed format and to submit the Performance Security within a specified time so as to become a Supplier.
- 2.13. NPPA:** The National Pharmaceutical Pricing Authority, the Government regulatory agency that controls the price of pharmaceutical formulations in India.
- 2.14. Period of Contract:** the rate price mentioned for the drugs by the Bidder shall be valid for a period of one year from the date execution of Contract.
- 2.15. Price Bid** shall have the meaning as ascribed to it in Clause 3.5 of this Tender Document.
- 2.16. Purchase Order** means an order issued by the Purchaser to the Supplier informing to supply the required quantity of the drugs at the contract price and to supply to various consignees as mentioned in the purchase order.
- 2.17. Revised Manual of Office Procedures for Procurement by Government Departments:** Government Resolution no. Bha.kha.sa-2014/ Pra. Kra. 82/Section-III/Industry-4, dated 1.12.2016 issued by Department of Industries and Labour, Government of Maharashtra and any subsequent amendments/revisions thereto.
- 2.18. Risk Purchase** is the additional cost incurred by the Purchaser in making alternate purchases of the quantity defaulted by the Supplier from other sources at a higher cost as compared to Price Bid quote.
- 2.19. Supplier:** is the selected Bidder(s) to whom Purchase Order(s) is placed on fulfilling the qualification criteria and terms and conditions laid down in this Tender Document.

2.20. Supply Schedule means the schedule for supply of drug which shall be adhered to for supply as per Clause 4.10 unless altered with mutual consent on the basis of the movement /consumption of drugs, exigencies and other reasons suiting the requirements of Purchaser.

2.21. Tender Document: The document published by the Purchaser containing the details of the drugs/medicines/consumables to be purchased, the quantity and delivery, and which includes designs, specifications, quality requirements and other Specific/General conditions which will govern the contract on acceptance of the Bid.

2.22. Unit: means the smallest unit of the drug(s) for which rates are to be quoted and to be made available on demand. The rate to be given on the price bid shall be quoted for this basic unit as mentioned in the BOQ.

Note: The words and expressions used in this Tender Document, but not defined, shall have the same meaning as respectively assigned to them under the prevailing Applicable Laws.

Section 3: General Terms and Conditions

This section deals with the general conditions of contract and contains the following terms & conditions governing the tender.

3.1. Responsibility of verification of contents of Tender Document

- 3.1.1. It shall be the responsibility of the Bidders to read/examine all instructions, forms, terms and specifications in the Tender Document and confirm that the required documents as specified in Clause 3.5 are duly uploaded. Failure to furnish any information required by the Purchaser in any respect shall result in the rejection of bids, without any notice.

3.2. Authorized Signatory for the Tender Document

- 3.2.1. Only authorized signatory identified and nominated in power of attorney submitted in the format prescribed in Annexure 7, shall be eligible to sign all documents and annexure related to the Tender Document. It is advisable for the Bidder to authorize only that person for this Tender Document, who is salaried employee of the Bidder. Further, the Bid shall be typed or written in indelible ink and the authorized signatory of the Bidder shall alone digitally sign and upload all required documents and annexures. All the alterations, omissions, additions or any other amendments made to the Bid shall be initialed by the person(s) signing the Bid. The Bid shall contain page numbers.

3.3. Period of Validity of Bid

- 3.3.1. The bid shall remain valid for a period of 180 days from the Bid Due Date (envelope no.1). Prior to the expiration of the bid validity the Purchaser may request the Bidders to extend the bid validity for the period as required by the Purchaser.

3.4. Earnest Money Deposit (EMD)

- 3.4.1. A Bidder needs to furnish/pay a minimum EMD amount of Rs.2.00 lacs, however if the Bidder is submitting its quote for more than 8 drugs, then for each additional drug beyond 8 number of drugs, the EMD amount shall correspondingly increase by a value of Rs. 25,000/- per additional drug, subject to a maximum of Rs. 5.00 lacs, unless exempted under Clause 3.4.7. In case, the value of EMD submitted by the Bidder does not correspond in value as per terms above, to the number of drugs for which the Price Bid is submitted, then the number of drugs starting at the top which corresponds to the actual EMD amount will only be considered for evaluation. For avoidance of doubt, the quoted items by the Bidder will be counted in sequence up to the value of EMD deposited. However, without minimum EMD the Bid will not be considered at all.
- 3.4.2. The payment of Earnest Money Deposit shall be made only online through payment gateway as provided in the tender portal and any other form such as Cheque/Cash/Postal order will not be accepted. The bids submitted without EMD will be summarily rejected.
- 3.4.3. Unsuccessful Bidder's EMD will be discharged/ returned within 2 months through e-Tender Portal after acceptance of NOA by the successful Bidder.
- 3.4.4. The Bidder shall not be entitled for any interest on EMD.

- 3.4.5. The successful Bidder's EMD will be discharged after signing the Contract and submitting the Performance Security & processing fee as stipulated.
- 3.4.6. Without prejudice to any other right or remedy that may be available to the Purchaser under the Tender Documents and/ or under the Contract, or otherwise, the Purchaser may forfeit the EMD and/or blacklist the Bidder upon occurrence any of the below-mentioned circumstances:
- i. A Bidder quotes prices higher than allowed as per DPCO, NPPA or higher than MRP; or
 - ii. a Bidder engages in an Unethical Practice as defined in Clause 3.21 of this Tender Document; or
 - iii. a Bidder withdraws its Bid during the period of Bid validity as specified in this Tender Document and as extended by mutual consent of the respective Bidder(s) and the Purchaser; or
 - iv. the Selected Bidder fails within the specified time limit -:
 - a. to sign and return the duplicate copy of NOA; or
 - b. To sign the Contract in accordance with terms and conditions or.
 - c. To furnish Performance Security and processing fee within the period prescribed therefor in this Tender Document.
 - v. The entire EMD paid shall be forfeited from the Bidder(s) whose manufacturing facilities were rejected on the grounds of non-compliance to statutory requirements as per the Applicable Laws.
- 3.4.7. Exemption for payment of EMD as per the Revised Manual of Office Procedures for Procurement by Government Departments (including Appendix-8 - Tender Form Fee and Earnest Money Deposits).

Micro and small-scale manufacturing industries registered under Micro, Small and Medium Enterprises development act 2006 are exempted from paying Tender Form Fees and EMD.

The above exemption is subject to submission of copy of Udyam Registration Certificate' or any other valid registration Certificate/Proof notified by the Government of India in respect of the drugs manufactured and quoted by them for participation in this tender floated by Purchaser. Further, the Purchaser reserves the right to inspect the manufacturing unit, whenever it is deemed necessary by it, in order to satisfy themselves with regard to verifying the credentials of the Bidder with respect to quality and production capacity and other relevant factors.

3.5. Submission of Bids

- 3.5.1. The Bidder shall submit the Bid no later than the date and time specified as the Bid Due Date, on the E-Tender Portal specified by the Purchaser, duly signed in digital form by the authorised signatory of the Bidder, by uploading the complete and legible scanned/digital copies of the Technical and Price Bids in BOQ/digital format (i.e. scanned copy of original signed documents and the supporting documents). The documents submitted in the Bid should be scanned in at least 100 dpi. It is further clarified, that if any document submitted with the Bid is not legible, the same may

not be considered by the Purchaser for further evaluation and the Bidder shall be solely responsible for any consequences thereof. The documents should be uploaded in the respective fields/space as per the provisions made on the portal. Each file of the document should be separate, and it should be uploaded in the given space/field only.

- 3.5.2. The Bid is to be submitted in accordance with the document downloaded including corrigendum issued thereto from the E-Tender Portal or the Official Website. The Bidder shall be responsible for its accuracy and correctness as per the version uploaded by the Purchaser and shall ensure that there are no changes caused in the content of the downloaded document.

To prepare and submit the bid/offer online all Bidders are required to have e-token based DIGITAL SIGNATURE CERTIFICATE. The Digital signature certificate should be obtained from competent authority;

- 3.5.3. **Technical Bid (Envelope No. 1):** The documents comprising the Technical Bid shall include:

1. Proof of Tender fees and EMD paid (if exempted as per Clause 3.4.7, attested copy of valid registration made by manufacturer for offered product under Micro & Small, Medium Industries Development Act, 2006).
2. Attested photocopy of valid manufacturing drug license with product list duly approved by the central licensing authority/ state licencing authority for each and every product quoted as per technical specification in the bid. The license must have been duly renewed up to date and the items quoted shall be clearly highlighted in the license. If quoted item is manufactured at different places, Manufacturing License & Performance certificate from all such places from respective Licensing Authority/ State Drug Authority should be enclosed. **However, Loan Licensee/ third party licensee are not allowed.**
3. World Health Organization - Good Manufacturing Practice certificate (with product list or COPP) highlighting the quoted products.
4. Copy of permission from DCGI for “New drug & Fixed Dose Combination”
5. Market Standing Certificate valid for a period of 3 years (i.e., for financial year 2022-23 and 2023-24 and 2024-25) as a manufacturer/importer for each drug(s) quoted in the tender within the Bid Due date. The period of Market Standing will be reckoned from the date of issue of Product permission. In case of an importer, their principal manufacturer located overseas should have 3 years market standing in India and the importer shall have 3 years market standing in the pharmaceutical field. In cases involving new drugs/ drugs out of patent period it is sufficient to possess relevant market standing certificate, as applicable.
6. Appendix-A (Schedule of Requirements)
7. Appendix-B (Checklist)
8. Annexure-1 (Technical Specifications and Compliance)
9. Annexure-2 (Letter Comprising Technical Bid)
10. Annexure-3 (Proforma for Production and Sale Statement)

11. Annexure-4 (Details of Manufacturing Unit)
12. Annexure-5 (Details of Items Quoted with Drug Code)
13. Annexure-6 (Annual Turnover Statement for Three Years) along with Copies of Balance Sheet and Profit and Loss Accounts for last three years i.e., (2022-23, 2023-24, 2024-25) certified by the Statutory Auditor or Chartered Accountant.
14. GST Registration certificate along with copy of the GST return of last quarter.
15. Annexure-7 (Format of Power of Attorney for signing of Bid) except for proprietorship.
16. Annexure-8 (Affidavit for Blacklisting)
17. Annexure-9 (Litigation Affidavit)
18. Annexure-10 (Mandate Form)
19. Incorporation / Registration Certificate of Bidder

Other Documents:

20. Schedule-1 (Contract Form)
21. Schedule-2 (Performance Security form)
22. Schedule-3 (Supply Schedule)
23. Schedule-4 (Schedule for Packing of Drugs and Medicines)
24. Schedule-5 (Bar Code & Advance Shipment Notification details)
25. Authorization letter nominating a responsible person of the Bidder to attend the meetings like pre-bid & negotiation meeting.

Note: In case the annual accounts for the latest financial year are not audited and therefore the Bidder cannot make it available, the Bidder shall give an undertaking to this effect and the statutory auditor/chartered accountant shall certify the same. In such a case, the Bidder shall provide the audited annual reports for 3 (three) years preceding the year for which the audited annual report is not being provided.

3.5.4. Bids submitted by special messenger, fax, telex, telegram, e-mail, or in any way other than on the specified e-tender platform for bidding, shall not be entertained and shall be rejected.

3.5.5. **Price Bid (envelope no. 2):**

- i. The composition and strength of each drug should be as per specifications given in Appendix A. Any variation, if found, will result in rejection of the tender/drug. However, the imported/combination drugs are allowed to be quoted in trade / brand name subject to clarification from the Bidder about composition of the drug.
- ii. The Price Bids of only those firms who qualify in the technical evaluation as per the terms herein, alone will be eligible for opening and evaluation of their Price Bid. Every Bidder shall submit their rates in the prescribed Proforma 'Price Bid Form' (BOQ) (refer Annexure 11) attached to the online bid document in Indian Rupees only for each of the required medicines separately on door delivery basis according to the unit in which prices has been sought. The Price Bid shall be submitted only online in the format given. The price bid (BOQ) file shall be

available to be downloaded from the E-Tender Portal and the Bidder shall quote the prices for respective drug as per Annexure 11 and upload the same on the E-Tender Portal. The Bidders shall not rename the BOQ files downloaded. Bidders are allowed to enter the Bidder's name & values only. Price bid in Annexure-11 should not be submitted in Technical Bid. If the Price Bid is submitted as part of the Technical Bid, the Bid will be rejected.

- iii. Bid for the supply of drugs, medicines, etc. with conditions like 'At Current Market Rates' shall not be accepted. The Purchaser shall not be responsible for damages, handling, clearing, transport charges and the same will not be paid. The deliveries should be made as stipulated in the Purchase Order placed with successful Bidder. Conditional bids are not accepted and liable for rejection.
- iv. The price shall be quoted on unit mentioned in BOQ format and not in respect of any other supply units. Any corrections in future in any respect shall not be entertained.
- v. The price quoted by the Bidder shall not in any case, exceed the controlled price, if any, fixed by the Central Government under DPCO OR NPPA and the Maximum Retail Price (MRP). The Purchaser will exercise the right to revise the price at any stage so as to conform with the controlled price or MRP as the case may be. The right under this clause will be exercised without prejudice to any other action that may be taken against the Bidder. Only landed cost (including all charges and taxes) mentioned in the Price Bid (quoted by the Bidder) is considered for rate comparison. Payment of all applicable taxes to concerned authority is the responsibility of the Bidder.
- vi. If at any time during the Period of Contract, the price of quoted items is reduced or brought down by any Applicable Law or by the Bidder itself, the Bidder shall be morally and statutorily bound to inform the Purchaser immediately about such reduction in the contracted prices. The Purchaser shall be empowered to reduce the rates accordingly.
- vii. In case of any enhancement in GST/other taxes due to statutory Act of the Govt. or any other taxes newly levied by Govt. after the date of submission of Bid and during the Period of Contract, the quantum of additional GST/other taxes so levied will be allowed to be charged extra as separate item without any change in price structure of the drugs approved under the bid. For claiming the additional cost on account of the increase in GST/other taxes, the Bidder should produce a letter from the concerned competent authorities for having paid additional GST/other taxes on the goods supplied to the Purchaser and can also claim the same in the invoice. Similarly, in case of reduction in GST / other taxes, the Purchaser will pay the reduced the GST / other taxes amount from the effective date of reduction in the aforesaid GST / other taxes.
- viii. **Fall Clause:** During the Period of Contract, the price at which Supplier will supply drugs shall not exceed the lowest price charged by Supplier to any other customer during the Period of Contract and in the event of Supplier supplies drugs to any other customer at a price below the price bid of Supplier, it shall

promptly furnish such information to Purchaser in order to correspondingly revise the rate of subsequent supplies.

- ix. The rates accepted by the Purchaser shall be binding on the Bidder during validity of the Bid and after execution of Contract for at least one year from the date execution of Contract. Any increase in the price will not be entertained till the completion of the Period of Contract.
- x. Purchases may be made on staggered basis as per the requirement of the Purchaser.
- xi. The total amount including taxes in column 8 of Annexure 11 (total amount in words) will only be considered for bid ranking.
- xii. In case it is noticed that a drug name has appeared in Annexure A for more than one time, then in such cases the Bidder having quoted the least price among such item will be considered as L1

3.6. Language

- 3.6.1. The Bid and all related correspondence and documents in relation to the bidding process shall be in English language. Supporting documents and printed literature furnished by the Bidder with the Bid may be in any other language provided that they are accompanied by translations of all the pertinent passages in the English language, duly authenticated and certified by the Bidder. Supporting materials, which are not translated into English, may not be considered. For the purpose of interpretation and evaluation of the Bid, the English language translation shall prevail.

3.7. Format of Bid

- 3.7.1. The Bidder shall provide all the information sought under this RFP. The Purchaser will evaluate only those Bids that are received in the required formats, in specified sequence, duly paginated and complete in all respects. Incomplete and /or conditional Bids shall be liable to rejection.

3.8. Number of Bids and Cost of bidding:

- 3.8.1. No Bidder shall submit more than one Bid under the Tender Document. A Bidder applying shall not be entitled to submit another Bid.
- 3.8.2. The Bidder shall bear all costs associated with the preparation and submission of their online bids and the Purchaser will in no case be responsible or liable for those costs, regardless of the conduct or outcome of the bidding process.

3.9. Amendment of Tender document:

- 3.9.1. At any time prior to the Bid Due Date, the Purchaser may amend the Tender Document by issuing addendum/corrigendum.
- 3.9.2. Any addendum/corrigendum as well as clarification thus issued shall be a part of the Tender Document. And it will be assumed that the information contained in the amendment will have been taken into account by the Bidder. Any addendum / corrigendum thus issued hereunder shall be hosted on the E-Tender portal.

- 3.9.3. To give prospective Bidders reasonable time in which to take the amendment into account in preparing their bids, the Purchaser shall extend, at its discretion, the deadline for submission of Bids, in which case, the Purchaser will notify all Bidders by placing it on Official Websites.

3.10. Acknowledgement by Bidder

- 3.10.1. It shall be deemed that by submitting the Bid, the Bidder has:

- i. made a complete and careful examination of the Tender Documents.
- ii. received all relevant information requested from the Purchaser.
- iii. satisfied itself about all matters, things, and information necessary and required for submitting an informed Bid, . of the Contract/ Purchase Order in accordance with the Tender Document and performance of all of its obligations thereunder.
- iv. acknowledged and agreed that inadequacy, lack of completeness or incorrectness of information provided in the Tender Documents shall not be a basis for any claim for compensation, damages, extension of time for performance of its obligations, loss of profits etc. from the Purchaser, or a ground for termination of the Contract by the Supplier;
- v. acknowledged that it does not have a Conflict of Interest; and
- vi. agreed to be bound by the undertakings provided by it under and in terms hereof.

- 3.10.2. The Purchaser shall not be liable for any omission, mistake or error in respect of any of the above or on account of any matter or thing arising out of or concerning or relating to the tender process or the bidding process, including any error or mistake therein or in any information or data given by the Purchaser.

3.11. Right to accept or reject any or all Bids.

- 3.11.1. Notwithstanding anything contained in this Tender Documents, the Purchaser reserves the right to accept or reject any Bid and to annul the bidding process and reject all Bids, at any time without any liability or any obligation for such acceptance, rejection, or annulment, and without assigning any reasons therefor. In the event that the Purchaser rejects or annuls all the Bids, it may, in its discretion, invite all eligible Bidders to submit fresh Bids hereunder.

- 3.11.2. The Purchaser reserves the right to reject any Bid if:

- i. at any time, a material misrepresentation is made or uncovered, or
- ii. the Bidder does not provide, within the time specified by the Purchaser, the supplemental information sought by the Purchaser for evaluation of the Bid.

Such misrepresentation/ improper response shall lead to the disqualification of the Bidder.

- 3.11.3. If disqualification/ rejection of a Bidder occurs after the Bids have been opened and the lowest Bidder gets disqualified/ rejected, then the Purchaser reserves the right to:

- i. invite the remaining Bidders to match the Lowest Bidder rate in accordance with the Tender Documents; or
- ii. take any such measure as may be deemed fit in the sole discretion of the Purchaser, including annulment of the bidding process.

- 3.11.4. In case it is found during the evaluation or at any time before signing of the Contract or after its execution and during the period of subsistence of Purchase Order that one

or more of the qualification conditions have not been met by the Bidder, or the Bidder has made material misrepresentation or has given any materially incorrect or false information, the Bidder shall be disqualified forthwith if not yet appointed as the Supplier either by issue of the NOA or entering into the Contract, and if the Bidder has already been issued the NOA or has entered into the Contract, as the case may be, the same shall, notwithstanding anything to the contrary contained therein or in this Tender Document, be liable to be terminated, by a communication in writing by the Purchaser to the Bidder, without the Purchaser being liable in any manner whatsoever to the Bidder. The Purchaser shall be entitled to forfeit and appropriate the EMD or Performance Security, as the case may be, as damages, and without prejudice to any other right or remedy which the Purchaser may have under this Tender Document, the Contract, Purchase Order or otherwise.

- 3.11.5. The Purchaser reserves the right to verify all statements, information and documents submitted by the Bidder in response to the Tender Documents and the Bidder shall, when so required by the Purchaser, make available all such information, evidence and documents as may be necessary for such verification. Any such verification or lack of such verification by the Purchaser shall not relieve the Bidder of its obligations or liabilities hereunder nor will it affect any rights of the Purchaser thereunder.
- 3.11.6. The Purchaser may, in its sole discretion and on grounds of reciprocity, disqualify a Bidder, if any or all of its constituents are entities incorporated in a country where an entity incorporated in India does not have similar rights of bidding for contracts contemplated hereunder.

3.12. Pre-Bid Meeting

- 3.12.1. The pre-bid meeting will be held at the date, time and venue mentioned in the bid notice and Bid Schedule. The Bidder shall note that, any corrigendum issued regarding this bid notice will be published on the <https://mahatenders.gov.in>.
- 3.12.2. A prospective Bidder requiring any clarification of the Tender document shall contact the Purchaser by letter or email to submit their suggestions/ observations/ Queries if any, preferably prior to the date of pre bid meeting on GEM portal or through Maha Tender Portal.
- 3.12.3. Only suggestions / observations related to Tender Documents, received in writing within stipulated time will be discussed and clarified in pre-bid meeting and any modification of the Tender documents, which may become necessary as a result of pre-bid meeting, shall be made by Purchaser exclusively through the issue of an addendum/ corrigendum. The bid uploaded shall be read along with any modification. Authorized representatives of prospective Bidder(s) can attend the said meeting and obtain clarification regarding specifications, scope of works & tender conditions. Authorized representatives should have authorization letter to attend the pre-bid meeting, subject to the condition that queries are submitted in time.
- 3.12.4. Any amendment to the Tender Document shall be placed on the E-Tender Portal (<https://mahatenders.gov.in>)/ GEM portal or on official website- mmgpa.maharashtra.gov.in
- 3.12.5. The Bidder will not be communicated separately regarding the amendment.

- 3.12.6. Non-attendance at pre-bid meeting shall not be a cause for disqualification of the Bidder. The suggestions/ objections/ queries received in pre-bid meeting may not be considered, if the same are not in consonance with the requirement of the bid. Purchaser reserves the right to reject the same.

3.13. Clarifications post Pre bid Meeting

- 3.13.1. Bidders requiring any clarification on the Tender Document may notify the Purchaser in accordance with Clause 3.12. They should send in their queries on or before the date specified in the Bid schedule of bidding process. The Purchaser shall endeavour to respond to the queries within reasonable time. The Purchaser will post all the queries and its responses on the E-Tender Portal (<https://mahatenders.gov.in/>) GEM portal or on official website- mmgpa.maharashtra.gov.in
- 3.13.2. The Purchaser may respond to the questions raised or clarifications sought by the Bidders in writing. However, the Purchaser reserves the right not to respond to any question or provide any clarification, in its sole discretion, and nothing in this Clause shall be taken or read as compelling or requiring the Purchaser to respond to any question or to provide any clarification.

3.14. Modification/substitution/withdrawal of Bids

- 3.14.1. No Bid shall be modified, substituted or withdrawn by the Bidder on or after the closing time on the Bid Due Date.
- 3.14.2. Any alteration/ modification in the Bid or additional information or material supplied subsequent to the closing time on the Bid Due Date, unless the same has been expressly sought for by the Purchaser, shall be disregarded.

3.15. Proprietary data

- 3.15.1. All documents and other information supplied by the Purchaser or submitted by a Bidder to the Purchaser shall remain or become the property of the Purchaser. Bidders are to treat all information as strictly confidential and shall not use it for any purpose other than for preparation and submission of their Bid. The Purchaser will not return any Bid, or any information provided along therewith.

3.16. Correspondence with the Bidder

- 3.16.1. No tenderer shall contact MMGPA, on any matter relating to its Bid, from the time of bid opening to the time of contract is awarded.
- 3.16.2. Any effort by the tenderer to influence MMGPA, in the purchasers bid evaluation, bid comparison or contract award decisions may result in rejection of the tenderers bid.
- 3.16.3. The tenderer shall not make any attempt to establish unsolicited and unauthorized contact with MMGPA or tender scrutiny committee after opening of the bids and prior to the notification of award and any attempt by any tenderer to bring to bear extraneous pressure on the tender accepting authority, tender inviting authority or tender scrutiny committee, shall be sufficient reason to disqualify the tenderer.
- 3.16.4. Notwithstanding anything contained in clause (3.16.3.) above MMGPA, may seek bonafide clarifications from the Bidders relating to the bid submitted by them during the evaluation of bids.

3.17. Opening and Evaluation of Bids

- 3.17.1. The Purchaser shall open the Technical Bids on the Bid Due Date as specified in Bid Schedule, online portal www.mahatenders.gov.in/ GEM Portal and in the presence of the Bidders who choose to attend.
- 3.17.2. The Purchaser will subsequently examine and evaluate Bids in accordance with the provisions set out in this Tender Document.
- 3.17.3. Bids of firms who have furnished all the required documents for each of the drug quoted alone will be considered. A firm quoting for more than one drug and if the required/proper document is not furnished for any of the drug(s), then offer of that drug(s) will be rejected. Utmost care should be taken to see that all the required/proper documents are uploaded.

3.17.4. Test of Responsiveness:

Prior to evaluation of Bids, the Purchaser shall determine whether each Technical Bid is responsive to the requirements of the Tender Document. A Technical Bid shall be considered responsive if:

- i. it is received as per the specified format and sequence mentioned in Clause 3.5;
- ii. the following mandatory documents are submitted with the Bid online, as on Bid Due Date:
 - a. in case of exemption claimed for Tender Fee and EMD, attested copy of valid registration made by manufacturer for offered product under Micro & Small, Medium Industries Development Act, 2006.
 - b. Copy of valid manufacturing drug license with product list duly approved by the Licensing Authority/ State drug authority for each and every product quoted as per specification in the bid. The license must have been duly renewed up to date and the items quoted shall be clearly highlighted in the product permission with their respective drug code as mentioned in the Appendix-A;
 - c. World Health Organization-Good Manufacturing Practice (with product list or COPP);
 - d. it is accompanied by the Technical Specifications and Compliance as per the Annexure-1;
 - e. it is accompanied by the Letter comprising Technical Bid as per the Annexure-2;
 - f. it is accompanied by the Proforma for Production and Sale Statement as per the Annexure-3;
 - g. it is accompanied by the Details of Manufacturing Unit as per the Annexure-4;
 - h. it is accompanied by the Annual Turnover Statement for Three Years) along with Copies of Balance Sheet and Profit and Loss Accounts for last three years i.e., (2022-23, 2023-24, 2024-25) certified by the Statutory Auditor or Chartered Accountant., as per the Annexure-5;

- i. it is accompanied by the Copy of certificate of incorporation/registration along with charter documents and GST Registration certificate;
 - j. it is accompanied by the Affidavit for Blacklisting as per theAnnexure-8;
 - k. it is accompanied by the Affidavit for Litigation as per theAnnexure-9; and
- iii. it is not non-responsive in terms hereof.

The Purchaser reserves the right to reject any Bid which is non-responsive and no request for alteration, modification, substitution or withdrawal shall be entertained by the Purchaser in respect of such Bid. Provided, however, that the Purchaser may, in its discretion, allow the Bidder to rectify any infirmities or omissions if the same do not constitute a material modification of the Bid.

- 3.17.5. The technical evaluation shall be on the basis of documents submitted and relevant standards of pharmacopoeia and provisions of Drugs and Cosmetics Act 1940 and Drugs and Cosmetics Rules 1945. Each item/medicine will be evaluated separately. Purchaser may at its discretion call for any documents for verification and the Bidder shall be duty bound to produce the same documents before the tender evaluation committee within stipulated time period. The status of bidders/items after technical bid evaluation will be published on the portal www.mahatenders.gov.in/GEM Portal and shall be final.
- 3.17.6. After the evaluation of Technical Bids, the Purchaser would announce a list of qualified Bidders who will be eligible for opening of their Price Bids. All communications relating to qualification shall be uploaded on Official Website. The Purchaser will not entertain any query or clarification from Bidders who fail to qualify.
- 3.17.7. Any information contained in the Bid shall not in any way be construed as binding on the Purchaser, its agents, successors or assigns, but shall be binding against the Bidder if the Contract is subsequently awarded to it on the basis of such information.
- 3.17.8. The Purchaser reserves the right not to proceed with the bidding process at any time without notice or liability and to reject any or all Bid(s) without assigning any reasons.

3.18. Confidentiality

- 3.18.1. Information relating to the examination, clarification, evaluation, and recommendation of the Bidders shall not be disclosed to any person who is not officially concerned with the process or is not a retained professional advisor advising the Purchaser in relation to, or matters arising out of, or concerning the bidding process. The Purchaser will treat all information, submitted as part of Bid, in confidence and will require all those who have access to such material to treat the same in confidence. The Purchaser may not divulge any such information unless it is directed to do so by any statutory entity that has the power under law to require its disclosure or is to enforce or assert any right or privilege of the statutory entity and/or the Purchaser or as may be required by law or in connection with any legal process.

3.19. Clarifications regarding Evaluation

- 3.19.1. To facilitate evaluation of Bids, the Purchaser may, at its sole discretion, seek clarifications from any Bidder regarding its Bid. Such clarification(s) shall be provided within the time specified by the Purchaser for this purpose. Any request for clarification(s) and all clarification(s) in response thereto shall be in writing.
- 3.19.2. If a Bidder does not provide clarifications sought under Clause 3.19.1 above within the prescribed time, its Bid may be rejected. In case the Bid is not rejected, the Purchaser may proceed to evaluate the Bid by construing the particulars requiring clarification to the best of its understanding, and the Bidder shall be barred from subsequently questioning such interpretation of the Purchaser.
- 3.19.3. Bidder shall ensure that, all correspondence with the Purchaser shall be through the official email id mentioned in Annexure 4 submitted by the Bidder.

3.20. Selection of Bidder

- 3.20.1. The Bidders are required to register on the e-Tender Portal for submission of their Bids in accordance with the procedure set out therein. Bidders are requested to visit the e-Tender Portal for the details related to online registration and submission of Bids. A Bidder may familiarize itself with the e-Tender Portal and in accordance with the instructions given on the e-Tender Portal (Bidders Manual Kit) and the terms of the Tender Document, submit its Bid. To participate in the bidding process, the Bidder should complete all stages of purchase, download of Tender Document from e-Tender Portal and undertake the final Bid submission through the e-Tender Portal. Bids which are submitted on the e-Tender Portal alone will be accepted by the Purchaser .
- 3.20.2. A Bidder may submit its Price Bid for one or more drugs in accordance with terms of this Tender Document. A Bidder is required to furnish all the specified documents in respect of each drug for which the Bidder submits its Price Bid.
- 3.20.3. Bids of Bidder who have furnished all the required documents in respect of each of the drug quoted alone will be considered. If a Bidder does not submit the required document complete in all respects as per the terms herein, then offer related to such drug(s) will be rejected. Utmost care should be taken to see that all the required documents are uploaded.
- 3.20.4. The Bidder's whose Bids are determined to be responsive to the requirements outlined in clause 3.17.4 shall be eligible for technical evaluation in accordance with clause 4.1 of the Tender Document.
- 3.20.5. The Bidder who meets the technical eligibility criteria and requirements of Supporting documentation (as per clause 4.1) shall be eligible for opening of the Price Bid.
- 3.20.6. After the conclusion of Price Bid opening (Envelope 2), the lowest offer of the Bidder(s) for the respective drug(s) will be considered for negotiation and respective L1 rates shall be arrived after negotiation for the drug(s). The Bidder(s) offering the L1 rate for the specified drug(s) will be declared as the selected Bidder for those drug(s) ("**Supplier**").
- 3.20.7. In the event that 2 (two) or more Bidders are qualified in terms hereof as L1 (referred to as "tie bidders"), then such Bidder having the higher production capacity as per

the eligibility criterion would be given first preference. Further, if tie Bidders are found to be having the same production capacity, then the Bidder having higher average annual turnover as per the eligibility criterion shall be taken into consideration and would be given first preference. Such Bidder shall execute necessary Contract as specified in the Tender Document. On depositing the required amount as Performance Security and on execution of the agreement, such Bidder will be eligible for the placement of Purchase Orders.

- 3.20.8. This tender is governed under the provisions of the Revised Manual of Office Procedures for Procurement by the Government Departments” issued by Department of Industries and Labour, government of Maharashtra vide GR dated 1.12.2016 or any subsequent amendment thereto. The purchase preference shall be in full adherence to the guidelines specified under the aforementioned manual. In case of any inconsistency between the purchase preference mentioned in this Tender Document, the same shall be resolved in accordance with provision of Revised Manual of Office Procedures for Procurement by the Government Departments and the same shall prevail.
- 3.20.9. Subject to clause 3.20.8 above, the Bidders from Maharashtra and / or MSME and / or SC/ST MSME, as the case may be, who have qualified for Price Bid (Envelope 2) opening, will be informed by the Purchaser of the lowest rate received for the drugs quoted by such Bidder and inviting their consent to match with the lowest rate for those drug(s). The Bidders who agree to match L1 Price Bid, will be considered as L1 Bidder. For the purpose of this Tender, the Bidders from Maharashtra would have the meaning as ascribed in Clause 3.1.2.8 of the Revised Manual of Office Procedures for Procurement by the Government Departments or any subsequent amendment thereto. On depositing the required amount as Performance Security and on execution of the Contract, such Bidder will be eligible for the placement of Purchase Orders for the quantities determined as per the provisions of “Revised Manual of Office Procedures for Procurement by the Government Departments.
- 3.20.10. The Bidder mentioned at clause 3.20.9 above, who matches the L1 Price Bid, on placement of Purchase Order, will be deemed as L1 Bidder for the purpose of the tender and all provisions of the Tender Documents applicable to L1 Bidder will apply mutatis mutandis to the Supplier who matches the L1 Price Bid.
- 3.20.11. Purchaser will issue Notification of Award to the L1 Bidder (L1) specifying the quantity for which the Tender is awarded and requiring the Bidder to execute a contract in the prescribed format and to furnish the Performance Security within 15 days from the issuance of Notification of Award, so as to become a Supplier.
- 3.20.12. The L1 bidder shall within 2 (two) days from the receipt of NOA submit to Purchaser its acceptance to Notification of Award.
- 3.20.13. The L1 Bidder on submission of acceptance to Notification of Award and Performance Security to the satisfaction of Purchaser, shall execute necessary Contract as per the format specified in Schedule-1 for the supply of the tendered quantity of such drug(s) as specified in the Tender Document.

3.21. Unethical Practices

- 3.21.1. The Purchaser as well as Bidders shall observe the highest standard of ethics during the procurement and execution of such contracts. For the purpose of this Tender Document Unethical Practices shall mean the following:
- i. “Corrupt practice” means the offering, giving, receiving or soliciting of anything of value to influence the action of a public official in the procurement process or in contract execution;
 - ii. “Fraudulent practice” means a misrepresentation or omission of facts in order to influence a procurement process or the execution of a Contract to the detriment of Purchaser and includes collusive practice among bidders (prior to or after bid submission) designed to establish bid prices at artificial non-competitive levels and to deprive the Purchaser of the benefits of free and open competition;
 - iii. “Collusive practice” means a scheme or arrangement between two or more bidders, with or without the knowledge of the Purchaser, designed to establish bid prices at artificial, non-competitive level; and
 - iv. “Coercive practice” means harming or threatening to harm, directly or indirectly, persons or their property to influence their participation in the procurement process or effect the execution of the Contract.
- 3.21.2. The Purchaser will reject a bid for award if it determines that the bidder recommended for award and/or execution of Contract has directly or through an agent has engaged in the abovementioned unethical practices at any point of time during bidding process or after execution of Contract. The Purchaser reserves the right to terminate the Contract, forfeit EMD/ Performance Security (as applicable) and debar/blacklist the Supplier from participating in any future Tender published by the Purchaser.

3.22. Code of Integrity

- 3.22.1. Any person participating in a procurement process shall-
- i. Not offer any bribe, reward or gift or any material benefit either directly or indirectly in exchange for an unfair advantage in procurement process or to otherwise influence the procurement process.
 - ii. Not misrepresent or omit or mislead or attempts to mislead so as to obtain a financial or other benefit or avoid an obligation.
 - iii. Not indulge in any collusion, bid rigging or anti-competitive behavior to impair the transparency, fairness and progress of the procurement process.
 - iv. Not misuse any information shared between the Purchaser and the Bidders with an intent to gain unfair advantage in the procurement process.
 - v. Not obstruct any investigation or audit of a procurement process.
 - vi. Disclose conflict of interest, if any; and
 - vii. Disclose any previous transgressions with any Entity in India or any other country during the last three years or any debarment by any other procuring entity.

3.22.2. The Purchaser will reject a bid for award if it determines that the Bidder recommended for award and/or execution of Contract has directly or through an agent has breached the above Code of Integrity at any point of time during bidding process or after execution of Contract. The Purchaser reserves the right to terminate the Contract, forfeit EMD/ Performance Security (as applicable) and debar/blacklist the Supplier from participating in any future Tender published by the Purchaser.

3.23. Payment Provisions

3.23.1. Payment against supply order issued under this bid will be made by Chief Account & Finance officer, Maharashtra Medical Goods Procurement Authority, Mumbai. 100% Payment shall be made on fulfilling the below mentioned conditions:

- i. upon submission of following documents by the Supplier:
 - a. 3 copies of Supplier's invoice,
 - b. Receipt and acceptance certificates issued by the consignees on E-Aushadhi, and
 - c. Batch wise in-house Lab Report.
- ii. Purchaser shall make payments of the quantity supplied for drugs which are declared to be of standard quality (SQ) by Purchaser empaneled labs.

3.23.2. Payments towards the supply of Drugs will be made strictly as per the rules of Purchaser. The payment will be made through RTGS/ NEFT. The Bidder shall furnish the relevant details in original to make the payment through RTGS/NEFT and the change of bank account during the validity of the bid will not be entertained normally.

3.23.3. The Purchaser shall have every right to deduct the pending dues on account of loss, compensation, or any remedial action in monetary terms from the said payment. The Supplier shall not agitate the said issue in future.

3.24. Termination of Contract

3.24.1. The Purchaser reserves the right to terminate the Contract on the below mentioned grounds:

- A. In case the Drugs are declared “Misbranded” ‘Adulterated’ & Spurious’ as per the applicable laws:
 - i. The contract of the Bidder for the said item will be cancelled.
 - ii. The extra expenditure incurred if any because of Risk Purchase shall be recovered from the Supplier.
 - iii. EMD or Performance Security, as applicable, of the Supplier will be forfeited.
 - iv. Purchase cost of full order if paid, irrespective of its consumed quantity shall be recovered from the Supplier from the outstanding bills or Performance Security .
 - v. The drugs which are not used but belong to the said substandard batch shall be destroyed by the concerned DDO in the presence of/or under intimation to Food and Drug Administration officials. The necessary expenditure incurred for this shall be recovered from the Supplier.
 - vi. The Supplier will be debarred from participating in any future Tender published by the Purchaser as per the terms of this Tender Document.

- B. In case the Drugs are declared “Not of Standard Quality” as per the Applicable Laws:
- I. The cancellation of Contract for the specified item shall be decided by the Chief Executive Officer, Maharashtra Medical Goods Procurement Authority, Mumbai, after reviewing the severity of sub-standard quality of item with the FDA Maharashtra. The testing report issued by FDA approved laboratory regarding quality shall be final & binding on the Supplier.
 - II. The extra expenditure incurred if any because of Risk Purchase shall be recovered from the Supplier.
 - III. Purchase cost, if paid, of full order irrespective of its consumed quantity shall be recovered from the Supplier from the outstanding bills or Performance Security.
 - IV. The goods which are not used but belong to the said substandard batch shall be destroyed by the concerned DDO in the presence of/or under intimation to Food and Drug Administration officials. The necessary expenditure incurred for this shall be recovered from the Supplier.
- C. In case it is determined, post award of Tender, that the Supplier has charged prices higher than allowed as per DPCO, NPPA for the quoted drug or higher than MRP (only in cases where DPCO, NPPA rates are not available), or has failed to supply the drugs consistently, the Supplier will be declared as fraudulent and defaulter and in such case: -
- I. The extra expenditure incurred due to higher prices charged over and above ceiling price as per DPCO, NPPA / MRP (as applicable) or in case of Risk Purchase shall be recovered from the Supplier.
 - II. The Contract shall be terminated, Supplier’s EMD/Performance Security, as applicable, will be forfeited and the Supplier will be debarred for next three years from participating in any future Tender published by the Purchaser.
- D. In case if found that the Bidder has submitted forged documents the following actions will be taken against the Bidder: -
- I. A police case will be filed against the bidder.
 - II. The Bidder’s EMD or Performance Security, as applicable, will be forfeited.
 - III. The bidder will be debarred for next three years from participating in any future Tender published by the Purchaser.
 - IV. The Contract already entered into will be liable for termination.
- E. In case if found that the drugs & other item supplied by the Supplier have been declared “Not of Standard Quality” as per the Applicable Law the actions will be taken as per Maharashtra Medical Goods Procurement Authority Regulations, 2025 and terms outlined in this Tender Document.
- F. In case, the Supplier becomes bankrupt or otherwise insolvent, the Purchaser reserves the right to terminate the contract at any time, by serving written notice to the Supplier without any compensation, whatsoever, to the Supplier, subject to

further condition that such termination will not prejudice or affect the rights and remedies which have accrued and/or will accrue thereafter to the Purchaser.

G. Termination for convenience: - The Purchaser reserves the right to terminate the contract, in whole or in part for its (Purchaser's) convenience, by serving written notice on the Supplier at any time during the currency of the Contract. The notice shall specify that the termination is for the convenience of the Purchaser. The notice shall also indicate inter alia, the extent to which the Supplier's performance under the Contract is terminated, and the date with effect from which such termination will become effective.

The Purchaser will be at liberty to terminate the contract either wholly or in part on 30-day notice. The Supplier will not be entitled for any compensation whatsoever in respect of such termination.

3.25. Dispute Resolution

- 3.25.1. In the event of any question, dispute or differences in respect of Contract or terms and conditions of the Contract or interpretation of the terms and conditions or part of the terms and conditions of the Contract arises, the parties shall make every effort to resolve, amicably by direct informal negotiation.
- 3.25.2. In the event of failure to settle the dispute amicably between the parties, the same shall be referred to the sole arbitrator as mutually appointed by the parties. In case, the parties fail to appoint the sole arbitrator mutually, in such case, board of three arbitrators, of whom each Party shall appoint one, and the third arbitrator shall be appointed by the two arbitrators so selected, and in the event of disagreement between the two arbitrators, the appointment shall be made in accordance with the Indian Arbitration and Conciliation Act, 1996 and the rules made thereunder.
- 3.25.3. The arbitrators shall make a reasoned award (the "Award"). Any Award made in any arbitration held pursuant to this Clause 3.25 shall be final and binding on the parties as from the date it is made, and the Supplier and the Purchaser agree and undertake to carry out such Award without delay.
- 3.25.4. The arbitration proceedings shall be carried out as per the Indian Arbitration and Conciliation Act, 1996 and the rules made thereunder.

3.26. Governing law and jurisdiction

- 3.26.1. This Contract shall be construed and interpreted in accordance with and governed by the laws of India, and the courts at Mumbai shall have exclusive jurisdiction over matters arising out of or relating to this Contract.

3.27. Indemnification

- 3.27.1. The Supplier shall indemnify the Purchaser against all actions, suit, claims and demand or in respect of anything done or omitted to be done by Supplier in connection with the Contract and against any losses or damages to the Purchaser in consequence of any action or suit being brought against the Supplier for anything done or omitted to be done by the Supplier in the execution of the Contract. .

3.28. Saving clause:

3.28.1. No suits, prosecution or any legal proceedings shall lie against the Purchaser, or any person for anything that is done in good faith or intended to be done in pursuance of this tender.

3.29. Force Majeure

3.29.1. For purpose of this clause, Force Majeure means an event beyond the control of successful Purchaser and not involving the successful tenderers fault or negligence and which is not foreseeable and not brought about at the instance of, the party claiming to be affected by such event and which has caused the non-performance or delay in performance. Such events may include, but are not restricted to, acts of the Purchaser either in its sovereign or contractual capacity, wars or revolutions, hostility, acts of public enemy, civil commotion, sabotage, fires, floods, explosions, epidemics, quadrant quarantine restrictions, strikes excluding by its employees, lockouts excluding by its management, and freight embargoes. Scarcity of raw materials/increase in the cost of raw material/shifting/up gradation of manufacturing facilities and power cut are not considered as Force Majeure.

3.29.2. If a Force Majeure situation arises, the successful tenderer shall promptly notify the Purchaser in writing of such conditions and the cause thereof with satisfactory documentary proof, within ten days of occurrence of such event. The time for making supply may be extended by the Purchaser as its discretion for such period as may be considered reasonable.

3.29.3. In case due to a Force Majeure event the MMGPA is unable to fulfil its contractual commitment and responsibility, the MMGPA will notify the successful tenderer accordingly and subsequent actions taken on similar lines described in above sub-paragraphs.

Section 4: Specific Terms and Conditions

This section deals with the specific conditions of contract and contains the following terms & conditions governing the tender.

4.1. Eligibility Criteria and Supporting Documents to be Submitted

Sr. No.	Basic Requirement	Specific Requirement	Documents required
1	Registered Legal Entity	The Bidder may be a natural person, private entity, government-owned entity, registered under applicable laws in India (“ Bidder ”). The Bidder shall be – a. Registered with the GST Authorities.	Copy of certificate of incorporation/registration along with charter documents like copy of Memorandum and Articles of Association, and other registration documents according to the nature of entity.

Sr. No.	Basic Requirement	Specific Requirement	Documents required
		b. Should have a valid PAN number.	Copy of GST Registration certificate issued by GSTN authorities. Copy of PAN Card.
2	Licenses	The Bidder must have valid manufacturing license/ import license for the drugs quoted as per technical specifications in the Tender Documents: a. Only manufacturer/ importer (i.e., authorized agent/subsidiary of the foreign manufacturer) will be allowed as Bidder. b. Loan Licensee / third party licensee are not permitted to participate under this Tender Document.	Self-attested copies of original manufacturing license in Form 25, 28, 28-D, 28-E, MD-5, MD-9 etc. approved by the Licensing authority along with retention Licenses. Or Self-attested copies of original import license in Form-10 or Form 10-A (as applicable), Form-41, MD-15 etc., and licenses in Form 20B/21B approved by the corresponding licensing authority with retention licenses. Or Self-attested copies of licenses in Form CT-20, Form CT-23 in case of new drugs. Further, during supply of drugs, if the drugs are manufactured at more than one premises, the Bidder shall submit the applicable license such as the manufacturing license, for each such premises mandatorily. Authority letter of the original manufacturer for importing the product for which bid is offered or agreement between foreign manufacturer and importer. The drugs quoted shall be highlighted in the product permission with their respective drug code as mentioned in the

Sr. No.	Basic Requirement	Specific Requirement	Documents required
			Appendix-A (Schedule of Requirement)
3	Certifications/ Registration	WHO-GMP (WHO - Good Manufacturing Practices Certificate) Certificate issued by the licensing authority. The WHO-GMP certificate must not be older than one year from the Bid Due Date in case where validity is not mentioned in the certificate. The WHO-GMP certificate of all the manufacturing plants, of which products have been quoted, should be submitted. The Importer should produce WHO-GMP / COPP of the manufacturing firm or a certificate which is at par with WHO-GMP issued by exporting countries like US-FDA approval, etc. In the case of imported drugs, labels and product literature of all quoted products must be submitted.	For manufacturers, WHO-GMP certificate with product list OR COPP, Quality management System (QMS) as per Medical Devices Rules, 2017 issued by the licensing authority wherever applicable. For importers, labels and product literature of all quoted drug(s) must be uploaded with WHO-GMP or certificate which is at par with WHO-GMP issued by the authorities of exporting countries of their principal manufacturing company or firm. Authority letter from OEM for the offered drug and valid import license issued by licensing authority. Import Export Certificate (IEC Code) for importer.
4	Past Experience	The Bidder must submit particulars of quantity of the past supplies of drugs made as per Annexure 3. Note: At least 25% quantity for similar drug as specified in the Annexure 1- Technical Specification and	Certificate issued by a statutory auditor/chartered accountant (as attached Annexure-3)

Sr. No.	Basic Requirement	Specific Requirement	Documents required
		in the Schedule of Requirements' must have been supplied for any of the last 3 (three) financial years preceding the Bid Due Date, i.e., 2022-23 ,2023-24,2024-25.	
5	Average Annual Turnover	Average Annual Turnover (in last three financial years 2022-23,2023-24,2024-25 shall be Rs 5,60,0000 /-	Certificate issued by a statutory auditor/chartered accountant (as attached Annexure-6) along with audited financial statements confirming the average annual Turnover of the Bidder during the stated financial years must be submitted. The turnover should be as certified by the chartered accountant/statutory auditor (specifying UDIN) and having valid registration. Note: In case the annual accounts for the latest financial year are not audited and therefore the Bidder cannot make it available, the Bidder shall give an undertaking to this effect and the chartered accountant/statutory auditor shall certify the same. In such a case, the Bidder shall provide the audited financial statements for the financial year preceding the latest financial year for which the audited annual report is not being provided.
6	Net Worth	The net worth of the bidder in the financial year immediately preceding the Bid Due Date should be positive .	Certificate issued by a statutory auditor/chartered accountant (as attached Annexure-6). The Net Worth should be as certified by the chartered accountant/statutory auditor (specifying UDIN) and having valid registration.

Sr. No.	Basic Requirement	Specific Requirement	Documents required
7	Production Capacity	Production capacity of the original drug manufacturer must be minimum 1.5 times of the quoted order quantity in last one financial year.	Certificate of Statutory Auditor/Chartered Accountant. As per Annexure 4.
8	Market Standing Certificate	Bidder should mandatorily possess 3 years Market Standing Certificate (i.e., for financial year 2022-23 and 2023-24 and 2024-25) as a manufacturer/importer for each drug(s) quoted in the tender as on Bid Due Date. The period of Market Standing will be reckoned from the date of issue of product/drug permission. In case of an importer, their principal manufacturer located overseas should have 3 years market standing in India and the importer shall have 3 years market standing in the pharmaceutical field. In cases involving new drugs/ drugs out of patent period it is sufficient to possess relevant market standing as applicable.	Market Standing Certificate as issued by Central or State Licensing Authority under the Applicable Law. Bill of landing of a foreign manufacturer and bill of entry of the importer, mentioning the country of origin. Notarized/ certified copy of Drug Controller General of India, new Delhi, for permission for Items coming under, “New Drug and Fixed Dose Combinations” in form 45/46 as per Drugs & Cosmetic Act and Rules. Relevant period Market standing certificate for new drug/drugs out of patent period issued by licensing authority.
9	Proof of distribution network	Bidders, if not having their registered office, manufacturing unit, or distribution facility within Maharashtra State, must have an established and operational distribution network in Maharashtra State at the time of bid submission.	Proof of distribution network— such as a valid drug selling/distribution license issued by the state’s licensing authority, along with details of authorized distributors (names, addresses, contact details) — must be submitted as part of the bid documentation Annexure 4 (A) and Copy of License of Sale / Distribution issued by State’s Licensing Authority

Sr. No.	Basic Requirement	Specific Requirement	Documents required
10	Blacklisting or banned	On the Bid Due Date, the Bidder should not be blacklisted or debarred by any ministry/department /attached offices/sub-ordinate offices under Government of India and any State government, Autonomous bodies (established by Central/State govt), any Central/State PSUs for unsatisfactory past performance, corrupt, fraudulent or any other unethical business practices. The bidder, whose drug has been declared as misbranded, spurious or adulterated or any criminal case in respect of the above is pending in any court, as on Bid Due Date shall not be eligible to participate for that particular drug, in the bid. Similarly, a Bidder convicted by court of law shall not be eligible to participate in the bid. The Bidder(s) quoting for this Tender Document should not have been convicted as on Bid Due Date by any court of law in India/overseas in lieu of deficiency noticed in the any of the quoted drug(s) in the tender and tender should not be submitted for such drugs for which conviction was pronounced by any court of law).	Affidavit as per Annexure 8 Non-Conviction Certificate issued by licensing authority/ State FDA

Sr. No.	Basic Requirement	Specific Requirement	Documents required
11	Litigation	The Bidder should not be involved in any major litigation such as fraud, FEMA violations and any other civil or criminal proceedings that may have an impact of affecting or compromising the delivery of services as required under this contract.	Affidavit as per Annexure 9.
12	EMD/Bid Security	The EMD indicated under Clause 3.4 unless exempted under Clause 3.4.7, shall be the Rs. 25,000/- per item of drug quoted subject to minimum of Rs.2.00 lacs and maximum of Rs. 5.00 lacs.	EMD payment shall be done as per the Clause 3.4.
	EMD Exemption	If a Bidder is a Micro and Small Enterprise (“MSEs”) / Small Scale Industry (“SSI”) then subject to submission of relevant documents as provided in this table, such Bidder may be exempted from submitting EMD in accordance with Appendix-8 of Revised Manual of Office Procedures for Procurement by Government Departments.	Requisite Certificate of Micro and Small-scale manufacturing industries registered under Micro, Small and Medium Enterprises Development Act, 2006.
13	Conflict of Interest	On the date of submission of the Bid, the Bidder should not have any conflict-of-interest situation.	Undertaking by the authorized signatory as per Annexure 2

4.1.1. **Conflict of Interest:** The Bidder participating in a bidding process must not have a Conflict of Interest. A Conflict of interest is considered to be a situation in which a party has interests that could improperly influence that party's performance of official duties or responsibilities, contractual obligations, or compliance with Applicable Laws and regulations.

A Bidder may be considered to be in Conflict of interest with one or more parties in bidding process if, including but not limited to:

- a. Have controlling partners/shareholders in common; or
 - b. Receive or have received any direct or indirect subsidy from any of them; or
 - c. Have the same legal representative for purposes of the Bid; or
 - d. 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 Bid of another Bidder, or influence the decisions of the Purchaser regarding the bidding process; or
 - e. The Bidder participates in more than one Bid in the same bidding process. Participation by a Bidder in more than one Bid for the same drug, will result in the disqualification of all Bids in which the Bidder is involved;
 - f. has a close business or family relationship with a staff with the Purchaser who: (i) are directly or indirectly involved in the preparation of the Tender document or specifications of the Tender Process, and/or the evaluation of bids; or (ii) would be involved in the implementation or supervision of resulting Contract unless the conflict stemming from such relationship has been resolved in a manner acceptable to the Purchaser throughout the tender process and execution of the Contract; and
 - g. The Bidder or any of its affiliates participated as a consultant in the preparation of the design or technical specification of the procurement of the drugs that are the subject of the Bid.
- 4.1.2. Bid should not be submitted for drug(s) for which the Bidder has been blacklisted/debarred either by Purchaser or by any other State/Central Government's organization/procurement agencies on the grounds of quality failure until completion of the penal period, and the bar subsists as on Bid Due Date.
- 4.1.3. Bid should not be submitted by any Bidder as a whole or for the specified drug(s) who have been blacklisted/debarred either by Purchaser or by any other State/Central Government's organization/procurement agencies/ autonomous bodies (established by Central/State govt), any Central/State PSUs on the grounds of unsatisfactory past performance, unethical practices such as fraudulent/corrupt practices etc., and the bar subsists as on Bid Due Date.
- 4.1.4. If any of the drug(s) of a Bidder have been blacklisted, during last 2 years from the Bid Due Date, such Bidder shall not be eligible to participate in this Tender Document for such drug(s). If it is found that the Bidder has quoted for any such drug as per the terms of this Tender Document, the Bidder shall be blacklisted for such particular drug for 2 (two) years and damages equivalent to EMD shall also be levied on the Bidder. In such situation, the Bid for remaining drugs (if quoted) will be considered further only if the damages are deposited before the completion of technical evaluation.
- 4.1.5. The Bidder, whose drug has been declared as of misbranded or spurious or adulterated quality and any criminal case is filed and pending in any court and subsists as on Bid Due Date, shall not be eligible to participate for that particular drug, under this Tender Document. Similarly convicted bidder shall also not be eligible to participate in the Bid.

- 4.1.6. If a Bidder has two or more separate manufacturing units at different sites/states, the Bidder will be allowed to submit only one Bid for all units but necessary document regarding separate manufacturing units will be submitted along with the Technical Bid. The Bidder will be allowed to submit only one Bid for one drug.
- 4.1.7. Any Bidder from a country which shares a land border with India will be eligible to Bid in this tender only if the Bidder is registered with the Competent Authority as provided in the Order (Public Procurement No. 1) dated 23rd July 2020 issued by the Ministry of Finance, Department of Expenditure Public Procurement Division. Provided further that the selected Bidder shall not be allowed to sub-contract works to any contractor from a country which shares a land border with India unless such contractor is registered with the Competent Authority as provided in the aforesaid Order. "Competent Authority" for the purpose of this clause means the Authority defined in Annex 1 of the Order (Public Procurement No. 1) dated 23rd July 2020 issued by the Ministry of Finance, Department of Expenditure Public Procurement Division.
- 4.1.8. This Tender Document is not transferable.
- 4.1.9. Any award of the Contract pursuant to this Tender Document shall be subject to the terms of Tender Documents.

4.2. Manufacturing/Importing and Product Permissions

- 4.2.1. Bidder should be a manufacturer duly licensed in Form 25/Form 28/ Form 28D/Form 28E/Form MD5 and Form MD9 (as per drugs quoted) with current validity/ retention issued by the State Licensing Authority (SLA)/Central Licensing Authority (CLA) as the case may be or a direct importer holding valid import license in Form 10 or Form 10-A as applicable, with Form 41/Form MD 15 issued by the Drugs Controller General of India (DCGI) accompanied with Licenses in Form 20B/21B with validity retention.
- 4.2.2. Bidder should have obtained permission to manufacture the drug(s) quoted strictly as per specification indicated in the Tender Document and in accordance with the standards specified in the Drugs and Cosmetics Act, 1940 from the competent authority such as state and/or central licencing authority.
- 4.2.3. In case of new drugs, import permission in Form CT-20 and manufacturing permission in Form CT-23 should be furnished in accordance with Rule 76/81 of the New Drugs and Clinical Trials Rules 2019 and Drugs and Cosmetics Act, 1940.
- 4.2.4. To ensure timely and smooth distribution of medicines, bidder, if not having their registered office, manufacturing unit, or distribution facility within Maharashtra State, must have an established and operational distribution network in Maharashtra State at the time of bid submission. Proof of such distribution network—such as a valid drug selling/distribution license issued by the state's licensing authority, along with details of authorized distributors (names, addresses, contact details) —must be submitted as part of the bid documentation

4.3. Minimum Tender Quantity

- 4.3.1. The details of the required drugs, etc., are shown in Appendix A. The Tender Quantity mentioned herein is not a fixed procurement quantity and it is only a tentative requirement and may be increased up to 50% or decreased by the Purchaser, at its discretion, depending on the actual need.
- 4.3.2. Though the tentative quantity is indicated in the Appendix A, the Purchaser, will confirm the actual requirement then and there only through Purchase Order(s). The Supplier shall supply the drugs only on the basis of the Purchase Order issued by the Purchaser. Any supply without a valid Purchase Order will not be accepted by MMGPA, for payment and the Purchaser, shall not be responsible for any loss on this account.
- 4.3.3. However, once the Purchase Order(s) are issued by the Purchaser, the Supplier should not renege from the commitment of supplying the quantity mentioned in the Purchase Order. The rates quoted shall also not be varied with the ordered quantity or the destination during the Contract Period.

4.4. Market Standing

- 4.4.1. Bidder should mandatorily possess 3 years Market Standing Certificate as a manufacturer/importer for each drug(s) quoted in the tender within the Bid Due date. The period of Market Standing will be reckoned from the date of issue of product/drug permission. In case of an importer, their principal manufacturer located overseas should have 3 years market standing in India and the importer shall have 3 years market standing in the pharmaceutical field. Also, the importer shall have due authorization for quoting drugs from the principal manufacturer along with relevant import licenses & marketing agreements as applicable. In the case of a new drug, bidder should possess relevant market standing as a manufacturer / importer from date of permission from DCGI and products (both of Plasma derived & recombinant categories) with USFDA certification, shall be considered with one-year global market standing. In cases involving any drugs out of patent period, it is sufficient to possess relevant market standing as applicable.
- 4.4.2. In cases of drug(s) with similar formulation but with varied strengths, market standing for 3 years for any strength of similar formulation shall be considered for all quoted drugs as equivalent, subject to possession of manufacturing license for the quoted drug(s) for a period not less than 3 years.
- 4.4.3. In case of imported drugs, market standing for the drug in international market would be considered for establishing eligibility regarding this particular clause of the bidding document. Also, if a bidder is manufacturing a drug abroad at various locations/countries and participating in the bid quoting a drug being manufactured at a particular place, market standing of the drug manufactured at other than particular place would be considered.
- 4.4.4. Bidder should have obtained permission to manufacture the drug(s) quoted as per specification in the tender and in accordance with the standards specified in the Drugs and Cosmetics Act, 1940 from the competent authority. The imported drug(s) should have valid import license by the competent authority. In both cases as indicated above, the permission provided by the Drug Controller General of India (DCGI) shall be in possession as applicable.

4.5. Inspection of Manufacturing Facilities

- 4.5.1. Purchaser may, at its discretion, conduct a inspection of the manufacturing premise.
- 4.5.2. Inspections of the production and related facilities of Bidders/ Suppliers will be at the discretion of the Purchaser. Such inspection may be at any stage after the Bidder is technically qualified for opening of Price Bid.
- 4.5.3. Where inspections are conducted as above, all parts of the manufacturing units including the quality control section will be subjected to rigorous inspection/auditing, irrespective of the items quoted. The Bidder/Supplier shall provide necessary co-operation for inspection of all the sections of the manufacturing unit. The denial of permission to inspect the manufacturing unit or failure to co-operate with the inspection of the different facilities or in providing information as per the details sought, will lead to disqualification.
- 4.5.4. The availability of plant & machinery, technical experts, analytical facilities of quality control lab etc., along with the compliance of WHO GMP regulations adopted for the production of quality assured products, all other parameters mentioned in the regulations shall be evaluated by the team for considering the eligibility of the firm. Claim of holding the valid certification/valid license will be of no avail for eligibility, if the procedures as stipulated in the standard operating procedures are not duly complied with, or if the available plant/ machinery are not in working condition at the time of inspection. Tender offer will be rejected/contract will be terminated with due notice in such cases.
- 4.5.5. Originals of all the documents uploaded/submitted in the Technical Bid as mentioned in Annexure-4 should be produced for verification during inspection. Failure to produce any of the original documents will result in the rejection of the tender offer deeming that the Supplier had made false statement at the time of the bid.
- 4.5.6. Key manufacturing areas may be photographed by the inspection team as a part of transparency and cross verification. Denial of permission for photographing may result in the rejection of Bid deeming that the Supplier had made false statement at the time of the Bid, if applicable, and/or the Purchaser may proceed with any actions available to it under the terms of this Tender Document.
- 4.5.7. Failure to observe any of the conditions of the licenses issued by competent authority, if reported by the inspection team will result in the rejection of the Bid deeming that the Bidder/Supplier had made false statement at the time of the Bid, if applicable, and/or the Purchaser may proceed with any actions available to it under the terms of this Tender Document.
- 4.5.8. The entire EMD or equivalent amount from the Performance Security shall be encashed, as the case maybe, paid by the Bidder(s) shall be forfeited whose manufacturing facilities were rejected on the grounds of non-compliance to statutory requirements. For the Bidder/Supplier claiming EMD exemption, the Bidder/Supplier shall be liable to pay damages to the Purchaser of an amount equivalent to the EMD for the drugs quoted by the said Bidder. In the event, the Bidder fails to pay the damages specified in this clause, the Purchaser reserves the

right to debar/blacklist the Supplier/Bidder for next three years from participating in any future Tender published by the Purchaser.

- 4.5.9. The Purchaser, or its authorized representative(s) shall have the right to inspect the factories of Bidders, before releasing any Purchase Order(s) or at any point of time after the Bid Due Date till the completion of the obligations as per the terms of this Tender Document/Contract, and also has the right to reject the Bid or terminate / cancel the Purchase Orders issued and/or not re-order, based on adverse reports brought out during such inspections.

4.6. Shelf Life of the Drugs

- 4.6.1. The supplied medicines and drugs should have the prescribed potency throughout the life period of drugs as prescribed in the Drugs and Cosmetics Act, 1940 (as amended) and rules there under and in relevant Pharmacopoeias. The “Life period of drugs” should be as per the Schedule P of the Drugs and Cosmetic Rules, 1945 (as amended).
- 4.6.2. Supplier(s) should supply the drugs to consignee within 30 days from the date of manufacture of that product. (Tablet, Capsule, Pessaries Ointment, Oral and Liquid preparation, Laboratory Chemicals, Kits, Injection and I.V Fluids etc.)

4.7. Method of Placing Purchase Orders

- 4.7.1. Subject to Clause 3.20, the following procedures will be adopted:
- i. After the conclusion of Price Bid opening (Envelope 2), the lowest offer of the Bidder for the respective drug(s) is considered.
 - ii. The Bidder, who has been declared as Supplier for respective drug(s), shall execute necessary Contract for the supply of the tendered quantity of such drug(s) as specified in the Tender Document. On depositing the required amount as Performance Security and on execution of the Contract, such Bidder(s) shall be eligible for the placement of Purchase Orders.

4.8. Execution of Contract

- 4.8.1. The lowest/matched Supplier shall execute a contract as per the form provided in Schedule-1 on a non-judicial stamp paper of value of as per The Maharashtra Stamp Act 1958 (stamp duty to be paid by the Supplier) within 15 days from the date issuance of the NOA from Purchaser. The cost of the stamp duty shall be borne by the Bidder. The Bidder shall not, at any time, assign, sub-let or make over the contract or the benefit thereof or any part thereof to any person or persons whatsoever.
- 4.8.2. All notices or communications relating to and arising out of this contract or any of the terms thereof shall be considered duly served on or given to the Bidder if delivered to him or left at the premises, places of business or abode as provided by the bidder.

4.9. Performance Security

- 4.9.1. The successful Bidder shall furnish Performance Security to the Purchaser at the time of execution of the Contract, for an amount of 3 % of the contract value for Bidders who are not MSE. In case the Bidder is MSE, the Bidders is required to provide Performance Security as per Clause 4.6 of Revised Manual of Office Procedures for Procurement of Goods by the Government Departments and any subsequent

amendment thereto. The Performance Security shall be valid for a period of 2 years from the date of signing of the Contract.

- 4.9.2. Performance Security can be in the form of ***Demand Draft or irrevocable Bank Guarantee in favour of the Maharashtra Medical Goods Procurement Authority, Mumbai from any Nationalized or Scheduled bank (Schedule-2).***
- 4.9.3. The Performance Security will be discharged by the Purchaser and returned to the Supplier not later than 60 days following the date of completion of the Supplier's performance obligations under the Contract.
- 4.9.4. The Performance Security shall be forfeited as a compensation for any loss resulting from the failure to perform the obligations under the Contract or in the event of termination of the Contract or in any event as the Purchaser thinks fit and proper, as the terms of this Tender Document.

4.10. Supply Conditions

- 4.10.1. Purchase Orders along with the place(s) of supply (consignee destinations) will be issued to the Supplier(s). The supplier should supply drugs within 30 days from the date of manufacturing at consignee destinations. No drugs will be accepted whose manufacturing date is beyond 30 days.
- 4.10.2. Within 2 days from the receipt of Purchase Orders, the Supplier(s) should provide the confirmation for the receipt with signed copy of Purchase Order and supply schedule as specified in Schedule-3 via email (maha.mmmpa2023@gmail.com).
- 4.10.3. The Supplier shall supply the entire ordered quantity within the timeline as specified in Maharashtra Medical Goods Procurement Authority Regulations 2025 dt: 21-03-2025, and subsequent amendments thereto. The supply period as per the said regulations is as below:

Category	Supply Period starting from the date of Purchase Order (in days)
Medicines and Medical supplies (Manufacturer)	45
Medicines and Medical supplies (Importer)	60
Essential Vaccines and highly specialized medicines	90

- 4.10.4. The Supplier shall supply the quantity specified in Purchase Order as per the timelines mentioned in the table above. If the Supplier fails to deliver any or all of the goods within the period(s) specified herein, the Purchaser shall, without prejudice to its other remedies under the Tender Document, shall recover from the Supplier as liquidated damages, a sum equivalent to 0.5% of the Contract price of the unsupplied drugs at the stipulated rate for each week or part thereof during which the delivery of such drugs may be delayed, as under:-
- i. In case of an order not exceeding Rs. 2.00 lakh in value –damage amount –at the rate of 0.5% per week (i.e., 0.0714% per day) subject to maximum limit of 10 %.

- ii. In Case of an order of Rs 2.00 lakh and above –damage amount –at the rate of 0.5% per week (i.e., 0.0714% per day) subject to maximum limit of 5 %.

- 4.10.5. If the Supplier fails to commence delivery as scheduled or to deliver the quantities ordered to him within the delivery period stipulated in the Contract, it shall be discretion of the Purchaser either (a) To extend the delivery period or (b) To cancel the Contract in whole or in part for the unsupplied quantities without any show cause notice.
- 4.10.6. All supplies will be scheduled for the period from the date of Purchase Order till the completion of the tender, as may be stipulated in the Purchase Order, subject to various conditions mentioned here under.
- 4.10.7. The Supplier(s) must submit an analysis report from a NABL accredited laboratory for every batch of drug along with invoice. In cases of Vaccines and Sera, the certificate of analysis from Central Drugs Laboratory, Kasauli shall be provided. In case of failure on part of the Supplier to furnish such report, the batch of drugs will be returned back to the Supplier and Supplier is bound to replenish the same with Govt. approved lab test report. The drugs supplied by the Supplier shall be of the best quality and shall comply with the specifications, stipulations and conditions specified in the tender.
- 4.10.8. Consignee shall not accept any shortages/damage in drug(s) at the time of receipt. It is Supplier's responsibility to fulfil or replace shortages/damage in drug(s) recorded at the time of receipt, as the case may be, within timelines as applicable as per the terms of the Purchase Order. Purchaser is not responsible for the excess stock of drug received, for which no order is placed.
- 4.10.9. If the Supplier fails to supply the drug(s) within the stipulated time, either fully or partly, Purchaser, is at liberty to place Purchase Orders either with other Bidders at the price offered by them or with alternate sources and in such cases the defaulted Supplier is liable to indemnify Purchaser, without any protest or demur, for the difference in cost incurred by Purchaser, and the Purchaser is entitled to recover the difference in cost from any amount due/payable to the defaulted Supplier.
- 4.10.10. Notwithstanding anything contained in Clause 4.10.9 above, the Supplier, after committing the default in supply either partly or fully, can inform the Purchaser, about its willingness to execute the Purchase Order. The Purchaser, at discretion, may consider the willingness of the Supplier on merit. However, such supplies will be subjected to the levy of liquidated damages and other penalties as stipulated in the Tender Document/ Contract and Purchase Order, at the discretion of the Purchaser.

4.11. Packing and Labeling

A. Packing

- 4.11.1. The drugs shall be supplied in the package specified in tender document and the packing and labelling shall be as mentioned in Schedule-4. Affixing of labels in smaller size will be treated as violation of tender conditions and fine will be deducted from the amount payable as per condition in Clause 4.12.2

- 4.11.2. 2D bar coding as per GS1 standard should be done on tertiary packing of the supplies as per the specifications given under Schedule-4.
- 4.11.3. 2D bar coding should be done on secondary packing for fixed variables as per the specifications given under Schedule-4.
- 4.11.4. The packing in each carton shall be strictly as per the specification mentioned in Schedule-4. Failure to comply with this shall lead to non-acceptance of the goods besides imposition of damages as per Clause 4.12 (ii). However, in case of poor / damaged packing, necessary replacement should be provided for damaged goods as notified by the Purchaser in such cases.
- 4.11.5. The caps of bottle preparations should not carry the name of the Supplier. The labels in the case of injectable preparations should clearly indicate whether the preparations are meant for Intravenous (IV), Intramuscular (IM), Intra Dermal (ID), Subcutaneous (SC) administration etc. The capsule shell should have the name of the drug.
- 4.11.6. It should be ensured that only first-hand fresh packaging material of uniform size, including bottle and vial, is used for packing. All primary packing containers should be strictly conforming to the specifications included in the relevant pharmacopoeia.
- 4.11.7. Packing should be able to prevent damage or deterioration during transit and storage. In the event of drugs supplied found to be not as per specifications in respect of their packing, the Purchaser, is at liberty to make alternative purchase of the drugs for which the Purchase Orders have been placed from any other sources or in the open market or from any other bidder who might have quoted higher rates, at the risk and the cost of the Supplier. In such cases the Purchaser, has every right to recover the cost and damages as mentioned in Clause 4.12.

B. Labelling

- 4.11.8. Labelling of the Drugs/Medical device shall be in strict compliance with Part-IX of the Drugs Rules, 1945/Chapter VI of Medical Device Rules, 2017 and other rules for the time being in force as approved by the appropriate Statutory authorities.

4.12. Schedule for packaging of essential drugs

(A) GENERAL SPECIFICATIONS:

- 1. No corrugated package should weigh more than 15 kilograms (i.e; product + inner carton + corrugated box).
- 2. All Corrugated boxes should be of “A” grade material i.e., Virgin quality.
- 3. All Essential Drugs should be packed only in first hand boxes.

FLUTE:

- 4. The corrugated boxes should be of narrow flute.

JOINT:

- 5. Every box should be preferably single joint and not more than two

joints.

STITCHING:

6. Every box should be stitched using pairs of metal pins with an interval of two inches between each pair. The boxes should be stitched and not joined using calico at the corners.

FLAP:

7. The flaps should uniformly meet and should not overlap each other. The flap when turned by 45 - 60° should not crack.

TAPE:

8. Every box should be sealed with Gum Tape/BoPP (Biaxially Oriented Polypropylene) tape running along the Top and lower opening.

CARRY STRAP:

9. Every box should be strapped with two parallel nylon carry straps (they should intersect).

LABEL:

10. Every corrugated box should carry a large outer label clearly indicating that the product is for "Maharashtra Govt. Supply - Not for Sale".

11. The product label on the carton should be large at least 15cms x 10cms dimension. It should carry the correct technical name, strength of the product, date of manufacturing, date of expiry, quantity packed and net weight of the box.

OTHERS:

12. No box should contain mixed products or mixed batches of the same product.

(B) SPECIFICATION FOR CORRUGATED BOXES HOLDING TABLETS /

CAPSULES / PESSARIES

1. The box should not weigh more than 7-8 Kilograms. The grammage of outer box should be 150 GSM and inside partition / lining should

be 120GSM.

2. The box should be of 5 ply with bursting strength of 9 Kg/ Cm²

(C) SPECIFICATIONS FOR OINTMENT / CREAM / GELS PACKED IN TUBES:

1. No corrugate box should weigh more than 7-8 Kgs.

2. Every Ointment tube should be individually packed in carton and then packed in 20's in a White board box, which may be packed in a corrugated box.

3. The grammage of outer box should be 150 GSM and inside partition / lining should be 120GSM.

(D) SPECIFICATIONS FOR INJECTABLE (IN VIALS AND AMPOULES):

1. Vials may be packed in corrugated boxes weighing upto 15 Kilograms. Ampoules should be packed in C.B weighing not more than 8 Kilograms.

2. Corrugated box for vials should be of 150 GSM (outer box should be 150 GSM and inside partition / lining should be 120 GSM) and 7 ply, while corrugated box for ampoules should be of 150 GSM (outer box should be 150 GSM and inside partition / lining should be 120 GSM) and 5 ply.

3. Bursting strength for CB boxes for

a. Vials: Note less than 13 Kg/Cm²

b. Amp: Note less than 9 Kg/Cm²

4. In the case of 10 ml Ampoules, 100 or 50 ampoules may be packed in a White board box. Multiples of White board boxes packed in corrugated box. In case of Ampoules larger than 10 ml, only 25 ampoules may be packed in a White board box with partition.

5. If the vial is packed in individual carton, there is no necessity for White board box packing. The individual carton may be packed as such in the corrugated box with center pad.

6. In case of ampoules, every White board box should carry 5 amps.

Cutters placed in a polythene bag.

7. Vials of Eye and Ear drops should be packed in an individual carton with a dispensing device. If the vial is of FFS/BFS technology, they should be packed in 50's in a White board box.

2D Bar Coding on Secondary Packing

DRUG CODE :
DRUG NAME :
BATCH NO :
SUPPLIER CODE:
SUPPLIER NAME:
MFG DATE:
EXPIRY DATE:



2D Bar Coding on Tertiary Packing

BOX NO:
PO NUMBER:
SUPPLIER CODE:
SUPPLIER NAME:
DRUG CODE:
DRUG NAME:
BATCH NO:
MFG DATE:
EXPIRY DATE:
BATCH QUANTITY:
INVOICE NO:



4.13. Aggregation and “parent-child” relationship serialisation

When the QR code or barcode on a **tertiary package** (like a shipping case or pallet) is scanned, the serial numbers of the **blister packs** inside (primary packaging) can be identified **without opening the package** due to a process called **aggregation** in pharma serialization.

- **Parent-Child Relationship:** During packaging, each blister (primary) receives a unique serial number and QR/DataMatrix code (the “child”). When these blisters are packed into cartons (secondary) and further into cases/pallets (tertiary), software systems link the serial numbers at each level by recording a “parent-child” relationship.
- **Central Database:** All this linking data (which blisters/serials are inside which carton, which cartons are inside which case) is uploaded to a secure database or the manufacturer's digital traceability system.
- **Tertiary Code Lookup:** When the tertiary-level code (unique QR code) is scanned, the system retrieves from the database the **complete list of serial numbers** for all individual blisters contained within that package—based on the links established during aggregation.
- **No Need to Unpack:** This allows supply chain operators or warehouses to quickly know exactly which blisters (their unique serial numbers) are inside any shipping case or pallet just by scanning the outside label.
- The identification is indirect: the tertiary code itself is not a “container” for all serial numbers, but is a reference that unlocks a digital “tree” of all linked codes for the inner packs.
- To ensure this facility the Bidder/Supplier shall ensure the aggregation process in packaging and provide the complete data of ‘Parent-Child’ linkage of the medical goods supplied and the access to its database to the same extent.

4.14. Deduction & Damages on Quality Failure

- 4.14.1. As soon as the drugs are received at the consignee location from the Suppliers, the details of the same shall be fed into the computer system (e-Aushadhi). Samples shall be drawn randomly from the Supplies from each batch. The samples drawn will be sent to the Quality Assurance (QA) cell of Purchaser.
- 4.14.2. After the samples are received at QA cell of Purchaser from randomly selected consignees, they will be segregated drug wise and batch number wise. The common batch of an item shall be mixed, and sample shall be drawn from pooled batch. Steps shall be taken to remove or hide the identity of the manufacturer and encode the formulations secretly. Encoded stickers shall be affixed on the blisters/strips/bottles/vials, as the case may be, so that the name and identity of the manufacturer gets hide but the contents of the sample are visible (Printed on the Sticker). The formulations / items assigned with secret unique codes will be sent to empaneled laboratories for analysis. Such sample should randomly be sent to laboratory. Within twenty-four hours of the receipt of samples, the information in appropriate format shall be uploaded by the empaneled laboratories on the e-Aushadhi portal. If any sample is received in a damaged condition by the laboratory, it shall not be analyzed, and the information thereof shall be sent immediately to the QA cell of Purchaser.
- 4.14.3. The empaneled laboratories shall analyze the drugs as per specifications and prescribed test protocols and submit its report to QA cell of Purchaser on e- Aushadhi

portal. The empaneled laboratory shall not disclose any of its information or report to anyone except the Purchaser. Every test report must have remark (i.e.) “Standard Quality” or “Not of Standard Quality”. Reports should have serial no., description of tests, specifications and results obtained as enumerated in relevant Acts and Rules. Reports shall be attached along with spectra/chromatography datasheets, if applicable.

- 4.14.4. If any sample sent to the empaneled laboratories fails in quality, the result may be confirmed with the other empaneled laboratories or Government Analyst before taking final decision.
- 4.14.5. If the drug fails in assay or any other parameters action shall be taken by the QA Cell immediately. The stock shall be frozen and kept separately until it is cleared. If the empaneled laboratories or government analyst confirm the failure of the drug to meet the standards, steps shall be taken to destroy or incinerate the stocks at the cost of the Supplier.
- 4.14.6. The report of the laboratory shall be communicated to the Supplier. The Supplier shall compensate for the substandard drug by supplying new batch of the same quantity within thirty days of the letter for information. If the Supplier fails to deliver the same within the stipulated timeline, then the Purchase Order for the item will be cancelled and no payment shall be made against that supply. For avoidance of doubt, it is clarified that the Damages shall be levied as per Clause 4.10.4 of this Tender Document.
- 4.14.7. If any of the drug/medicines supplied by the Supplier has been partially or wholly used or consumed after supply and are subsequently found to be in bad odour, unsound, inferior in quality or description or otherwise faulty or unfit for consumption, then the contract price for the quantity not consumed and informed to take back, will be recovered from the Supplier, if payment has already been made.
- 4.14.8. The Supplier shall furnish the source of procurement of raw material utilized in the formulations, if required by the Purchaser. The Purchaser reserves the right to cancel the Purchase Orders, if the source of supply is not furnished.
- 4.14.9. The decision of the Purchaser, or any officer authorized by it, as to the quality of the supplied drugs, medicines etc., shall be final and binding.
- 4.14.10. In the event of making alternative purchase, as specified in Clause 4.10 (ix) damage will be imposed on the Supplier. The excess expenditure over and above contracted prices incurred by Purchaser, in making such purchases from any other sources or in the open market or from any other bidder who has quoted higher rates and other losses sustained in the process, shall be recovered from the Supplier including through encashment of Performance Security or adjustment from any money payable to the Supplier under this Contract. Upon encashment and appropriation of the Performance Security, the Supplier shall, within [15 (fifteen)] days thereof, replenish, in case of partial appropriation, the Performance Security to its original level, and in case of appropriation of the entire Performance Security provide a fresh Performance Security, as the case may be, failing which the Purchaser shall be entitled to terminate the Contract. In all the above conditions, the decision of the Purchaser, shall be final and binding.

4.15. Blacklisting of Product/Tenderer on withdrawal of Tender:

- 4.15.1. If the Tenderer(s) fails to execute the agreement / to deposit performance security / to perform the obligations under the tender conditions / commits default in the performance of the contract, such Tenderers will be blacklisted for a period of 2 years by MMGPA., from the date of intimation besides forfeiture of Earnest Money Deposit (EMD) / Performance Security.
- 4.15.2. The Tenderers who have withdrawn after submitting their bids and before MMGPA finalizes and places the purchase orders either fully or partially, will be blacklisted for a period of 2 years from the date of intimation by MMGPA, apart from forfeiture of the Performance Security /EMD.

4.16. Blacklisting for Non-Supply of purchase Order(s):

- 4.16.1. If the supplier fails to execute at least 70% of the ordered quantity as mentioned in a single Purchase order and such part supply for any three purchase orders of the same drug, then the product of the supplier will be blacklisted and becomes ineligible to participate in any of the tenders for that particular drug(s) by MMGPA, for a period of 2 years from the date of intimation for blacklisting besides forfeiture of performance security of that product(s).
- 4.16.2. If the supplier supplies more than one drug and 50% of such drugs are blacklisted, the firm/company is liable to be blacklisted for a period of 2 years from the date of intimation besides forfeiture of performance security in full.
- 4.16.3. Purchase orders, if any, already issued before taking any blacklisting action or orders given in past will not be affected in view of action taken as per above guidelines but all strict quality checks shall be observed for each supply of products.
- 4.16.4. The blacklisting of particular product or company/firm will be done without prejudice to other penalties which may be imposed as per the conditions of Tender documents and also to other actions which may be initiated under Drugs and Cosmetics Act 1940 or any other law of MMGPA, will display names of such blacklisted product(s) and company/firm on its website and also circulate the same among other state Government / Central Government and its Drug procurement agencies including respective State Drugs Control Department where the company or firm is located.

4.17. Blacklisting of Product(s) based on Quality Test by the Empanelled Laboratories

- 4.17.1. Each and every batch of Drug/Medicines supplied by the supplier shall be subjected to quality test by the Empanelled laboratories.
- 4.17.2. The samples are collected from the Warehouses/consignees from each batch of supply of the same drugs and after eliminating the common batch, samples shall be taken in random, decoded and will be sent to the empanelled testing laboratories for testing the quality of drugs.
- 4.17.3. If such sample passes quality test in all aspects, MMGPA, will instruct its Warehouses/consignees to issue such drugs to various hospitals/Institutions for use.

- 4.17.4. Such quality passed batches if received after declaration of result of the earlier supply, the same will be again subjected to testing and the latest report of that particular batch will prevail upon the earlier results and binding on the entire quantity of the batch supplied and recovery will be made for the entire quantity of that batch irrespective of purchase order date or date of supply etc.
- 4.17.5. If the sample fails in quality test and report is received certifying that sample is not of standard quality, one more sample shall be drawn from the same batch and will be sent to Government Drugs Testing Laboratory for quality testing.
- i If such sample passes the quality test as per the report of Government Analyst, the drugs representing the sample shall be qualified for issue to various Institutions.
 - ii If such sample fails in the quality test, as per the report of the Government Analyst, the drugs of the batch are not qualified for issue and the supplier shall take back the drugs supplied in that batch, besides taking other actions as per the Tender conditions by MMGPA.
 - iii If such Sample fails in quality test for ASSAY content of less than 50% as per the Government Analyst report, such product of the tenderer will be blacklisted for two years beside forfeiture of Performance security of that product.
 - iv However, MMGPA, reserves the right to reject the drugs based on reports from empanelled laboratories with the applicable penal provisions.
- 4.17.6. If 3 batches of a particular drug supplied by the supplier is reported to be failing in Assay content (above 50% but below prescribed limit) / Sterility / BET / Toxicity / and or other parameters, then the particular drug of the firm shall be blacklisted and forfeiture of Performance Security of that particular product(s).
- 4.17.7. In all the cases the reports received from the Government Drug Testing Laboratory/ decision of MMGPA will be conclusive and final and binding on the suppliers.

4.18. Blacklisting of Product(s) based on Quality Test by Statutory Authorities

- 4.18.1. On intimation from Drugs Inspector(s) during their statutory sampling, that the particular drug has been reported to be of not of standard quality, the issue of available stock of the particular drug will be stopped. Further, the available stock of the product in hospitals will be retrieved. If the sample is reported to have less than 50% of content, the particular product will be blacklisted for 2 years from the date of intimation of blacklisting
- 4.18.2. If 3 batches of a particular drug supplied by the supplier is reported to be failing in assay content (above 50% but below prescribed limit) and/or other parameters, then the particular drug of the firm shall be blacklisted for a period of 2 years from the date of intimation.
- 4.18.3. If a single batch of any product(s) supplied by the company/firm declared as Adulterated/spurious/ Misbranded by the Government Authorities during the shelf life of the product supplied irrespective of tender period, the company/firm shall be blacklisted for a period of 5 years from the date of intimation.

4.19. Blacklisting of the Supplier on Quality Failure:

- 4.19.1. In case of any sample even in one batch, declared as Adulterated/spurious/Misbranded by the Government Authorities, the company/firm shall be blacklisted for a period of 5 years from the date of intimation besides forfeiture of performance security in full.
- 4.19.2. If the supplier supplied more than one drug and 50% of such drugs are blacklisted, the firm is liable to be blacklisted for a period of 2 years from the date of intimation.

4.20. Procedure for Blacklisting/Debarment:

- 4.20.1. On receipt of report from Govt. Analyst/Drug Testing Laboratory indicating that a particular Drug/Drugs is Not of Standard Quality/ Adulterated/ Spurious/ Misbranded (as the case may be), a show cause notice shall be issued to the supplier calling for explanation within 7 days from the date of notice. On receipt of explanation from the supplier, the CEO, MMGPA, may take appropriate action on merits of the case and impose penalty @ 25% of the value of the failed batch (or) 7.5% of the total supply value made in the particular purchase order (which ever higher) or blacklist the particular drug(s) of the company or firm as deemed fit besides forfeiture of Performance Security.
- 4.20.2. If a particular drug has been blacklisted according to the procedure stated above, the supplier is not eligible to participate in any of the tenders for that particular drug floated by the MMGPA, until the period of blacklisting is over.
- 4.20.3. 3If a supplier company/firm is blacklisted according to the procedure stated above, such supplier is not eligible to participate in any of the tenders floated by MMGPA until the period of blacklisting is over.

Appendix-A: Schedule of Requirements:

Sr. No.	Name of Drugs	Specification	Pack Size	Unit	Required Quantity	Required turnover
1	2 ACT (Adult) Strip	Total dose of Artesunate-600 mg divided over three days, SulphadoxinePyremethamine (1500 mg + 75 mg) single doseEach Combi Blister Pack containing 3 tablets of Artesunate (200mg each) and 2 tablets of Sulphadoxine Pyremethamine (750mg+37.5mg)First Row (Day 1) One tablet of Artesunate (200mg) and two tablets of sulphadoxinePyremethamine (750mg+ 37.5mg) Second Row(Day 2)- One tablet of Artesunate (200 mg), Third ROw(Day 3)-One tablet of Artesunate(200mg) & as per state Govt specifications	Combi Pack	Per Strip	1490	178800
2	3 ACT (0 -- 1years) Strip	Total dose of Artesunate-75 mg divided over three days, SulphadoxinePyremethamine (250 mg + 12.5mg) single doseEach Combi Blister Pack: containing 3 tablet of Artesunate (each tablet of Artesunate 25mg strength) and 1 tablet of SulphadoxinePyremethamine (250mg + 12.5mg First Row (Day-1) One Tab of Artesunate (25mg) and one tablet of Artesunate (25 mg) and one tablet of rtesunate (25 mg) and one tablet of Suphadoxine-Pyremethamine (250 mg + 12.5 mg) Second Row(Day 2)- One tablet of Artesunate (25 mg), Third ROw(Day 3)-One tablet of Artesunate(25mg)& as per state Govt specifications	Combi Pack	Per Strip	450	21600
3	4 ACT (1 -- 4 years) Strip	Total dose of Artesunate-150 mg divided over three days, SulphadoxinePyremethamine (500 mg + 25mg) single dose Each Combi Blister Pack containing 3 tablets of Artesunate (50mg each) and 1 tablet of SulphadoxinePyremethamine (500mg+25mg) First Row (Day 1)One tablet of Artesunate (50mg) and one tablet of sulphadoxinePyremethamine (500+25mg)Second Row(Day 2)- One tablet of Artesunate (50 mg), Third ROw(Day 3)-One tablet of Artesunate(50mg)& as per state Govt specifications	Combi Pack	Per Strip	450	25200

Sr. No.	Name of Drugs	Specification	Pack Size	Unit	Required Quantity	Required turnover
4	5 ACT (5 -- 8 years) Strip	Total dose of Artesunate-300 mg divided over three days, SulphadoxinePyremethamine (750 mg + 37.5mg) single dose Each Combi Blister Pack containing 3 tablets of Artesunate (100mg each) and 1 tablet of SulphadoxinePyremethamine (750mg+37.5mg) First Row (Day 1) One tablet of Artesunate (100mg) and one tablet of sulphadoxinePyremethamine (750mg+37.5mg) Second Row(Day 2)- One tablet of Artesunate (100 mg), Third Row(Day 3)-One tablet of Artesunate(100mg)& as per state Govt specifications	Combi Pack	Per Strip	460	40480
5	6 ACT (9 -- 14 years) Strip	Total dose of Artesunate-450 mg divided over three days,SulphadoxinePyremethamine (1000 mg + 50 mg) single doseEach Combi Blister Pack containing 3 tablets of Artesunate (150mg each) and 2 tablets of SulphadoxinePyremethamine (500mg+25mg) First Row (Day 1)One tablet of Artesunate (150mg) and two tablets of sulphadoxinePyremethamine (500mg+ 25mg) Second Row(Day 2)- One tablet of Artesunate (150 mg), Third Row(Day 3)-One tablet of Artesunate(150mg)& as per state Govt specifications	Combi Pack	Per Strip	670	69680
6	Activated Charcoal Oral	Activated charcoal/Magnesium hydroxide	500 gm Jar	Per Jar	76188	91426
7	Chloroquine base Tab 250 mg	Tab Chloroquine 250mg Aluminium blister strip of 10 tablets& as per state Govt specifications	10 Tab	Per Tab	143000	286000
8	Chloroquine Syrup 50 mg/5ml 60 ml	Each 5 ml contain ChloroquinePhasphate IP equivalent to chloroquine base 50 mg& as per state Govt specifications	60 ml Bottle	Per Bottle	500	8400
9	Chlorpheniramine Maleate Syrup 100 ml	Syrup Chlorpheniramine Maleate Each 5ml contain Pheniramine Maleate 2 mg - 100 ml	100ml Bottle	Per Bottle	1251890	11016632

Sr. No.	Name of Drugs	Specification	Pack Size	Unit	Required Quantity	Required turnover
10	Diazepam Oral Liquid 2mg/5ml	Diazepam Oral Liquid 2mg/5ml	30 ml Bottle	Per Bottle	44187	275727
11	Diethylcarbamazine citrate Tab I.P 100mg	Tab. Diethyl carbamazin 100 mg Aluminium Blister - Strip of 10 Tab	Strip of 10 Tablets	Per Tab	66990	26260
12	Enalapril Tab 2.5mg	Tab. Enalapril Maleate 2.5 mg	Strip of 10 Tablets	Per Tab	605188	135562
13	Ferrous fumarate Syrup 100ml Bottle	Sy. Ferrous fumarate 140 mg/ 5 ml, Bottle of 100ml	100ml Bottle	Per Bottle	524022	6791325
14	Fluconazole Tab 100mg	Fluconazole Tab 100mg	Strip of 10 Tablets	Per Tab	1740565	1740565
15	Gentamycin inj. 10mg	Inj. Gentamycin 10 mg/ml - 2 ml amp/Vial	2 ml	Per Amp/Vial	308639	513575
16	Human Insulin NPH/Isophane 40iu/ml	Highly purified bovine porcine isophane insulin susp 40 i.u./ml NPH insulin (or neutral protamine Hagedorn) (also known as Humulin N, Novolin N, Novolin NPH, NPH Iletin II, and isophane insulin),	10 ml Vial	Per Vial	26749	1797533
17	Ibuprofen Tab. 200mg	Tab. Ibuprofen 200 mg F/c Aluminium Blister - Strip of 10 Tablets	Strip of 10 Tablets	Per Tab	22960550	7347376
18	Isosorbide Dinitrate Tab 5mg	Tab. Isosorbide dinitrate 5 mg Aluminium Blister - Strip of 10 Tablets	Strip of 10 Tablets	Per Tab	407247	55386
19	Omeprazole Cap 40 mg	Omeprazole Cap 40 mg	Strip of 10 Tablets	Per Cap	202290	113282
20	Phenobarbitone Syrup 20mg/5ml, 60ml Bottle	Sy Phenobarbitone 20 mg per 5 ml - 60 ml Bottle	60 ml Bottle	Per Bottle	57742	1136363
21	Phenytoin Sodium ER Tablet 300 mg	Phenytoin Sodium ER Tablet 300 mg Aluminium Blister - Strip of 10 Tab	Strip of 10 Tablets	Per Tab	63844	182849
22	Primaquine Tab 15mg	Tab Primaquin 15 mg Aluminium blister strip of 10 tablets& as per state Govt specifications	Strip of 10 Tablets	Per Tab	259731	201082
23	Propranolol Tab 40mg	Tab.Propranalol HCL 40 mg	Strip of 10 Tablets	Per Tab	325536	106776
24	Pyridoxine Cap 50 mg	Cap Pyridoxine 50 mg	Strip of 10 Cap	Per Cap	1545000	988800
25	Quinine Inj 300 mg 2 ml Amp	Inj Quinine 300 mg 2 ml& as per state Govt specifications	2ml Amp	Per Amp	200	3200

Sr. No.	Name of Drugs	Specification	Pack Size	Unit	Required Quantity	Required turnover
26	Quinine Tab 300 mg	Tab Quinine Sulphate 300 mg Aluminium blister strip of 10 tablets & as per state Govt specifications	Strip of 10 Tablets	Per Tab	650	2028
27	Rifampicine Cap 600 mg	Cap Rifampicine 600 mg	Strip of 10 Cap	Per Cap	198000	1267200
28	Rifampicine Cap 450 mg	Cap Rifampicine 450 mg	Strip of 10 Cap	Per Cap	96500	552752
29	Rifampicine Cap 150 mg	Cap Rifampicine 150 mg	Strip of 10 Cap	Per Cap	70000	224000
30	Salbutamol 200mcg rotaCaps powder puff	Salbutamol 200 mcg per Cap - powder puff	Bottle of 30 Caps	Per Cap	168443	71420
31	Salbutamol Tab 2mg	Tab. Salbutamol Sulphate 2 mg Aluminium Blister - Strip of 10 Tab	Strip of 10 Tablets	Per Tab	1649816	197978
32	Senna Tablet, Granules, Powder	1 kg Packet Contains Senna Tablet, Granules, Powder	1 kg pkt	Per Packet	15000	1440000
33	Silver Amalgam	Each Capsule Contains Silver 48%, Copper 22% And Tin 30% 50 Capsule of silver alloy No Gamma 2 Amalgam With Mercury	1*50N Capsule Pack	Per Cap	1000	1200000
34	Thalidomine Tab 100 mg	Tab Thalidomine 100 mg	Strip of 10 Tablets	Per Tab	9500	342000
35	Topical Lignocaine (Xylocaine) 4%	Xylocaine 4% Topical 30 ml	30 ml Bottle	Per Bottle	35968	536643
36	Vitamin A Capsule 5000 IU	Cap. Vitamin A 5000 IU	Strip of 10 Cap	Per Cap	448992	172413

Delivery Terms : **To the consignee destination on door delivery basis
as per Tender conditions**

Appendix-B: Checklist

(Mandatory Documents to be uploaded online in the Technical Bid)

TECHNICAL BID DOCUMENTS.			Page No. (as per the Bid)
1	Proof of Tender fees and EMD paid (if exempted as per Clause 3.4, attested copy of valid registration made by manufacturer for offered product under Micro & Small, Medium Industries Development Act, 2006 (if applicable).	Yes / No	
2	Copy of certificate of incorporation/registration along with charter documents, and other registration documents according to the nature of entity	Yes / No	
3	Pan Card Details	Yes / No	
4	GST Registration certificate along with copy of the GST return of last quarter.	Yes / No	
5	Copy of manufacturing drug license with product list duly approved by the Licensing Authority/ State drug authority for each and every product quoted as per specification in the bid. The license must have been duly renewed up to date and the items quoted shall be clearly highlighted in the license	Yes / No	
6	World Health Organization-Good Manufacturing Practice certificate with either product list or COPP	Yes / No	
7	Copy of permission from DCGI for “New drug & Fixed Dose Combination”	Yes / No	
8	Market Standing Certificate as issued by Central or State Licensing Authority for relevant period against quoted Drugs	Yes / No	
9	Non-Conviction Certificate issued by Licensing Authority.		
10	Annexure-1 (Technical Specifications and Compliance)	Yes / No	
11	Annexure-2 (Letter Comprising Technical Bid)	Yes / No	
12	Annexure-3 (Proforma for Production and Sale Statement)	Yes / No	
13	Annexure-4 (Details of Manufacturing Unit)	Yes / No	
14	Annexure-5 (Details of Items Quoted with Drug Code)	Yes / No	

15	Annexure-6 (Annual Turnover Statement for Three Years) along with Copies of Balance Sheet and Profit and Loss Accounts for last three years i.e., (2022-23, 2023-24, 2024-25) certified by the Statutory Auditor or Chartered Accountant.	Yes / No	
16	GST Registration certificate along with copy of the GST return of last quarter.	Yes / No	
17	Annexure-7 (Format of Power of Attorney for signing of Bid) except for proprietorship	Yes / No	
18	Annexure-8 (Affidavit for Blacklisting)	Yes / No	
19	Annexure-9 (Litigation Affidavit)	Yes / No	
20	Annexure-10 (Mandate Form)	Yes / No	
21	Import Export Certificate (IEC Code) for Importer	Yes / No	

For the avoidance of any confusion, scanned copies of the above mentioned documents shall be uploaded online on the Official Website on or prior to the Bid Due Date.

Annexure 1: Technical Specifications and Compliance

Tender reference No: E-248/MMGPA/ Essential Medicines (2025-26)

Item Name: Following are the minimum requirements. Products offered must meet these parameters herein.

Sr. No	Item Name	Technical specification s/ composition of tender enquiry	Compliance on each parameter with detailed substantiation how the offered product meets the requirement.	If Column B=C (write Yes or No)	Generic Name / Brand Name (only for Importer)	Drug Mfg. License (Form 25 or 28 & 26)/ Medical devices/ Import License (Form 10 & 41)	MSM E/ SSI	Remarks, if any
	A	B	C	D	E	F	G	H
1	2 ACT (Adult) Strip	Total dose of Artesunate-600 mg divided over three days, SulphadoxinePyremethamine (1500 mg + 75 mg) single doseEach Combi Blister Pack containing 3 tablets of Artesunate (200mg each) and 2 tablets of Sulphadoxine Pyremethamine (750mg+37.5mg)First Row (Day 1) One tablet of Artesunate (200mg) and two tablets of sulphadoxinePyremethamine (750mg+ 37.5mg) Second Row(Day 2)- One tablet of Artesunate (200 mg), Third ROW(Day 3)-One tablet of Artesunate(200 mg) & as per state Govt specifications						

2	3 ACT (0 -- 1years) Strip	Total dose of Artesunate-75 mg divided over three days, SulphadoxinePyremethamine (250 mg + 12.5mg) single dose Each Combi Blister Pack: containing 3 tablet of Artesunate (each tablet of Artesunate 25mg strength) and 1 tablet of SulphadoxinePyremethamine (250mg + 12.5mg First Row (Day-1) One Tab of Artesunate (25mg) and one tablet of Artesunate (25 mg) and one tablet of Artesunate (25 mg) and one tablet of Sulphadoxine-Pyremethamine (250 mg + 12.5 mg) Second Row (Day 2)- One tablet of Artesunate (25 mg), Third Row (Day 3)-One tablet of Artesunate(25mg)& as per state Govt specifications						
3	4 ACT (1 -- 4 years) Strip	Total dose of Artesunate-150 mg divided over three days, SulphadoxinePyremethamine (500 mg + 25mg) single dose Each Combi Blister Pack containing						

		3 tablets of Artesunate (50mg each) and 1 tablet of SulphadoxinePyremethamine (500mg+25mg) First Row (Day 1) One tablet of Artesunate (50mg) and one tablet of sulphadoxinePyremethamine (500+25mg) Second Row (Day 2)- One tablet of Artesunate (50 mg), Third Row (Day 3)- One tablet of Artesunate (50mg) & as per state Govt specifications						
4	5 ACT (5 -- 8 years) Strip	Total dose of Artesunate-300 mg divided over three days, SulphadoxinePyremethamine (750 mg + 37.5mg) single dose Each Combi Blister Pack containing 3 tablets of Artesunate (100mg each) and 1 tablet of SulphadoxinePyremethamine (750mg+37.5mg) First Row (Day 1) One tablet of Artesunate (100mg) and one tablet of sulphadoxinePyremethamine (750mg+37.5mg) Second Row (Day 2)- One tablet of						

		Artesunate (100 mg), Third ROw(Day 3)-One tablet of Artesunate(100 mg)& as per state Govt specifications						
5	6 ACT (9 -- 14 years) Strip	Total dose of Artesunate-450 mg divided over three days,Sulphadox inePyremetham ine (1000 mg + 50 mg) single doseEach Combi Blister Pack containing 3 tablets of Artesunate (150mg each) and 2 tablets of SulphadoxineP yremethamine (500mg+25mg) First Row (Day 1)One tablet of Artesunate (150mg) and two tablets of sulphadoxinePy remethamine (500mg+ 25mg) Second Row(Day 2)- One tablet of Artesunate (150 mg), Third ROw(Day 3)-One tablet of Artesunate(150 mg)& as per state Govt specifications						
6	Activated Charcoal Oral	Activated charcoal/Magn esium hydroxide						

7	Chloroquine base Tab 250 mg	Tab Chloroquine 250mg Aluminium blister strip of 10 tablets & as per state Govt specifications						
8	Chloroquine Syrup 50 mg/5ml 60 ml	Each 5 ml contain Chloroquine Phosphate IP equivalent to chloroquine base 50 mg & as per state Govt specifications						
9	Chlorpheniramine Maleate Syrup 100 ml	Syrup Chlorpheniramine Maleate Each 5ml contain Pheniramine Maleate 2 mg - 100 ml						
10	Diazepam Oral Liquid 2mg/5ml	Diazepam Oral Liquid 2mg/5ml						
11	Diethylcarbamazine citrate Tab I.P 100mg	Tab. Diethylcarbamazine 100 mg Aluminium Blister - Strip of 10 Tab						
12	Enalapril Tab 2.5mg	Tab. Enalapril Maleate 2.5 mg						
13	Ferrous fumarate Syrup 100ml Bottle	Sy. Ferrous fumarate 140 mg/ 5 ml, Bottle of 100ml						
14	Fluconazole Tab 100mg	Fluconazole Tab 100mg						
15	Gentamycin inj. 10mg	Inj. Gentamycin 10 mg/ml - 2 ml amp/Vial						
16	Human Insulin NPH/Isophane 40iu/ml	Highly purified bovine porcine isophane insulin susp 40 i.u./ml NPH insulin (or neutral protamine						

		Hagedorn) (also known as Humulin N, Novolin N, Novolin NPH, NPH Iletin II, and isophane insulin),						
17	Ibuprofen Tab. 200mg	Tab. Ibuprofen 200 mg F/c Aluminium Blister - Strip of 10 Tablets						
18	Isosorbide Dinitrate Tab 5mg	Tab. Isosorbide dinitrate 5 mg Aluminium Blister - Strip of 10 Tablets						
19	Omeprazole Cap 40 mg	Omeprazole Cap 40 mg						
20	Phenobarbitone Syrup 20mg/5ml, 60ml Bottle	Sy Phenobarbitone 20 mg per 5 ml - 60 ml Bottle						
21	Phenytoin Sodium ER Tablet 300 mg	Phenytoin Sodium ER Tablet 300 mg Aluminium Blister - Strip of 10 Tab						
22	Primaquine Tab 15mg	Tab Primaquin 15 mg Aluminium blister strip of 10 tablets& as per state Govt specifications						
23	Propranolol Tab 40mg	Tab. Propranolol HCL 40 mg						
24	Pyridoxine Cap 50 mg	Cap Pyridoxine 50 mg						
25	Quinine Inj 300 mg 2 ml Amp	Inj Quinine 300 mg 2 ml& as per state Govt specifications						
26	Quinine Tab 300 mg	Tab Quinine Sulphate 300 mg Aluminium blister strip of 10 tablets& as per state Govt						

		specifications						
27	Rifampicine Cap 600 mg	Cap Rifampicine 600 mg						
28	Rifampicine Cap 450 mg	Cap Rifampicine 450 mg						
29	Rifampicine Cap 150 mg	Cap Rifampicine 150 mg						
30	Salbutamol 200mcg rotaCaps powder puff	Salbutamol 200 mcg per Cap - powder puff						
31	Salbutamol Tab 2mg	Tab.Salbutamol Sulphate 2 mg Aluminium Blister - Strip of 10 Tab						
32	Senna Tablet,Granules, Powder	1 kg Packet Contains Senna Tablet,Granules, Powder						
33	Silver Amalgam	Each Capsule Contains Silver 48%, Copper 22% And Tin 30% 50Capsule of silver alloy No Gamma 2 Amalgam With Mercury						
34	Thalidomine Tab 100 mg	Tab Thalidomine 100 mg						
35	Topical Lignocaine (Xylocaine) 4%	Xylocaine 4% Topical 30 ml						

Note:

- Pharmacopoeia standards IP/BP/USP etc. should be clearly mentioned against each drug/constituent of the formulation quoted as per the provisions of the Applicable Laws.
- Active ingredient used in formulation of item quoted shall be of mentioned Pharmacopoeia quality & Specifications.

Sign

Stamp

Date

MNGPA Tender

Annexure-2: Letter Comprising Technical Bid

**To,
Chief Executive Officer,
Maharashtra Medical Goods Procurement Authority,
1st Floor, Aarogya Bhawan,
Near CSMT Railway Station,
Mumbai 400001 (Maharashtra)**

Subject: Bid for the [***]**

Dear Sir,

With reference to your Tender document dated , I, having examined the Tender document and understood its contents, hereby submit my/our Bid for the aforesaid [*****]. The Bid is unconditional and unqualified.

1. I/ We acknowledge that the Purchaser will be relying on the information provided in the Bid and the documents accompanying such Bid for selection of the Supplier for the supply of the specified drugs, and we certify that all information provided therein is true and correct; nothing has been omitted which renders such information misleading; and all documents accompanying such Bid are true copies of their respective originals.
2. I/ We shall make available to the Purchaser any additional information it may find necessary or require supplementing or authenticate the Bid.
3. I/ We acknowledge the right of the Purchaser to reject our Bid without assigning any reason or otherwise and hereby waive, to the fullest extent permitted by applicable law, our right to challenge the same on any account whatsoever.
4. I/ We certify that in the last three years, we or our associates have neither failed to perform on any contract, as evidenced by imposition of a penalty by an arbitral or judicial authority or a judicial pronouncement or arbitration award, nor been expelled from any contract by any public authority nor have had any contract terminated by any public authority for breach on our part.
5. I/ We declare that:
 - (a) I/ We have examined and have no reservations to the Tender Document, including any Addendum / Corrigendum issued by the Purchaser.
 - (b) I/ We do not have any conflict of interest in accordance with the Tender Documents; and
 - (c) I/We have not directly or indirectly or through an agent engaged or indulged in any unethical practice, as defined in the Tender Document, in respect of any tender or request for proposal issued by or any agreement entered into with the Purchaser or any other public sector enterprise or any government, Central or State.
6. I/ We understand that you may cancel the bidding process at any time and that you are neither bound to accept any Bid that you may receive nor to invite the Bidders to Bid for

the Tender , without incurring any liability to the Bidders, in accordance with the provisions of the Tender Document.

7. I/ We believe that we satisfy(ies) the Annual Turnover and Net Worth criteria and meet(s) all the requirements as specified in the Tender Document and am/ are qualified to submit a Bid.
8. I/ We certify that in regard to matters other than security and integrity of the country, we or any of our associates have not been convicted by a Court of Law or indicted or adverse orders passed by a regulatory authority which could cast a doubt on our ability to undertake the Contract or which relates to a grave offence that outrages the moral sense of the community.
9. I/ We further certify that in regard to matters relating to security and integrity of the country, we or any of our associates have not been charge-sheeted by any agency of the Government or convicted by a Court of Law.
10. I/ We further certify that no investigation by a regulatory authority is pending either against us or against our associates or against our CEO or any of our directors/ managers/ employees / Partners /Trustees.
11. I/We further certify that we or any of our Associates are not barred by the Central Government/ State Government or any entity controlled by it, from participating in any supply of drugs contract, and no bar subsists as on the Bid Due Date .
12. I/ We undertake that in case due to any change in facts or circumstances during the bidding process, we are attracted by the provisions of disqualification in terms of the provisions of this Tender Document, we shall intimate the Purchaser of the same immediately.
13. I/ We hereby irrevocably waive any right or remedy which we may have at any stage at law or howsoever otherwise arising to challenge or question any decision taken by the Purchaser in connection with the selection of the Bidder, or in connection with the bidding process itself, in respect of the above-mentioned Contract and the terms and implementation thereof.
14. In the event of my/ our being declared as the Selected Bidder, I/we agree to enter into a Contract in accordance with the draft that has been provided to me prior to the Bid Due Date along with the Tender Document. We agree not to seek any changes in the aforesaid draft and agree to abide by the same.
15. I/ We have studied all the Bidding Documents carefully. We understand that except to the extent as expressly set forth in the Contract and Purchase Order, we shall have no claim, right or title arising out of any documents or information provided to us by the Purchaser or in respect of any matter arising out of or relating to the bidding process including the award of Contract.
16. The power of attorney for signing of Bid, as per format provided at Annexure-7 of the Tender Document, is also enclosed.
17. I/ We agree and undertake to abide by all the terms and conditions of the Tender Document.

18. I/ We offer a Tender Fee of INR [****] and EMD of INR [*****] to the Authority in accordance with the Tender Document.
19. I/We agree and understand that the Bid is subject to the provisions of the Tender Documents. In no case, I/We shall have any claim or right of whatsoever nature if the Contract is not awarded to me or our Bid is not opened or rejected.
20. The Price Bid has been quoted by me after taking into consideration all the terms and conditions stated in the Tender Document, our own estimates of costs and after a careful assessment of the all the conditions that may affect the price and implementation of the Contract.
21. I/We shall keep this offer valid for 180 days from the Bid Due Date as specified in the Tender Document.
22. I/ We hereby undertake to submit this Technical Bid for undertaking the aforesaid Contract in accordance with the Tender Documents and the Contract.

In witness thereof, I/we submit this Bid under and in accordance with the terms of the Tender Document.

Yours faithfully,

Date:

(Signature, Name and designation of the Authorised signatory)

Place:

(Name and seal of Bidder)

Annexure-3: Proforma for Production and Sale Statement

(For a period of last 3 Years preceding the Bid Due Date)

Name of the bidding entity:

Sr. No.	Year	Name and full Address of the Purchaser	Purchasing Entity (Gov./Semi Gov./Other)	Name of the Product	Purchase Order No. & Date	Purchase Order Quantity	Batch No.		PO Copy enclosed on Pg. No.
							Manufactured Qty	Sold Qty	
1	2022-23								
2	2023-24								
3	2024-25								

Add rows as per requirement.

Note:

1. In support of above statement, enclose the copies of supply orders and client's satisfactory certificates. All purchase orders should be enclosed in the serial as per the data provided in table above.
2. In case of importer, sold quantity shall be provided, production and manufactured good details may not be provided-
3. All the data of the bidding entity, as provided in the above table has been verified by undersigned Chartered Accountant/Statutory Auditor.
4. The issuer of this certificate must ensure that the above information/details are related to the bidding entity only.

Name, Membership number and signature of the Chartered Accountant/Statutory Auditor:

UDIN

Name and seal of the firm:

Location, Date:

Authorized Signature (*PoA holder*)

[In full and initials with Seal]:

Name and Title of Signatory:

Name of Bidder (Firm/ Organization's name):

Address:

Telephone:

Email:

(*Name and seal of the Bidder*)

[*Location, Date*]

Annexure-4: Details of Manufacturing Unit

(The details of manufacturing facility to be furnished)

Name of the Bidder and Office Address :

Factory Address :

PAN :

GST No. :

Phone Nos. :

Fax :

E-Mail¹ :

Date of Inception :

License No. & Date :

Issued by :

Valid up to :

Details of installed Production Capacity :

¹ Bidder shall ensure that, all correspondence with the Purchaser shall be through the official email id mentioned herein.

Details of Installed Production Capacity for 30 days
(In Terms of Unit Packs)

Tablets :

Capsules

General :

Beta-Lactum :

Injection

Ampoules :

Vials :

I.V.Fluids :

Sterile Powder:

Liquids

Suspension :

Syrups :

Drops :

Ointment :

Powders :

Antiseptics/
Disinfectants :

Name & designation of the authorized signatory:

Specimen signature of the authorized Signatory:

*The details of manufacturing unit should be for the premises where drugs quoted are actually manufactured

THE DETAILS OF FACTORY PREMISES

Person In-charge of Factory

Name :

Phone No. :

Mobile No. :

Nearest Landmark of Factory:

Layout

Km from Airport :

Name of the Airport and City:

Km from Railway Station :

Name of the Railway Station:

Km from Bus Stand :

Name of the Bus Stand:
And City

All the data provided in the above table has been verified by undersigned Chartered Accountant/Statutory Auditor.

Name, Membership number and signature of the Chartered Accountant/Statutory Auditor:

UDIN

Name and seal of the firm:

Location, Date:

Authorized Signature (*PoA holder*)

[In full and initials with Seal]:

Name and Title of Signatory:

Name of Bidder (Firm/ Organization's name):

Address:

Telephone:

Email:

(Name and seal of the Bidder)

[Location, Date]

Annexure-4 (A): Proof of Distribution Network

1. Sales / Distribution of License no :
(Copy of license is enclosed herewith)
2. Details of Authorized distributors in Maharashtra State :

Sr No	Name	Address	Contact Details
1			
2			

(Name and seal of the Bidder)

[Location, Date]

Annexure-5: Details of Items Quoted with Drug Code

1. Name of the firm:
2. Address as given in drug licence:
3. Drug Licence No. in Form 25 &28:
4. Import Licence No. :
5. Date of issue:
6. Validity:
7. Non-conviction Certificate obtained on:
8. Market standing certificate obtained on:
9. Details of endorsement for all products:

Sr. No.	Drug name	Specifications IP/BP/USP	Date of endorsement obtained from State Drugs Commissioner	Whether Endorsement is in Generic or Brand Name

Add As Many Rows as possible You Want to Add

(Additional column should be inserted asking date of permission from CDSCO, in case of all newly introduced drugs and Fixed dose combinations)

Annexure 6: Annual Turnover Statement for Three Years

(To be submitted on the letterhead of the Statutory Auditor/ Chartered Accountant of the Bidder)

M/s. is participating in tender No..... Items details as per Below Table.

Sr No of Item (as per Appendix A)	Name of Item	Required Turnover (In Rs)
Total Required Turnover		

The Average Annual Turnover and Net Worth details of M/s _____ for participation under the Tender Document are given below and certified that the statement is true and correct.

Sr. No.	Year	Turnover (In Rs.)
1	2022-23	
2	2023-24	
3	2024-25	
4	Average Annual Turnover of above 3 years	
Net Worth		
5	Net worth in the latest financial year preceding the Bid Due Date (positive/negative)	

“Net Worth” in case of Company shall mean the aggregate value of the paid-up share capital and all reserves created out of the profits and securities premium account, after deducting the aggregate value of the accumulated losses, deferred expenditure and miscellaneous expenditure not written off, as per the audited balance sheet, but does not include reserves created out of revaluation of assets, write-back of depreciation and amalgamation.

For other eligible entities, the Net Worth shall mean the amount derived by subtracting the liabilities from the assets as certified by the chartered accountant/statutory auditor having valid registration.

Note:

1. Certificate issued by a statutory auditor/chartered accountant along with audited financial statements confirming the average annual turnover of the Bidder during the stated financial years must be submitted on the letterhead of the statutory auditor/chartered accountant.
2. Provide supporting audited financial statements (Balance Sheets, Profit and Loss Statements, etc.) of the bidding entity.
3. The Net Worth of the bidder in the financial year immediately preceding the Bid Due Date should be positive.
4. “Turnover” for the purposes of this Tender Document shall mean the monetary value of

goods sold by the Bidder.

Name, Membership number and signature of the Chartered Accountant:

UDIN

Name and seal of the firm:

Location, Date:

Authorized Signature of Bidder (*PoA holder*)

[In full and initials with Seal]:

Name and Title of Signatory:

Name of Bidder (Firm/ Organization's name):

Address:

Telephone:

Email:

(*Name and seal of the Bidder*)

[*Location, Date*]

Annexure-7: Format for Power of Attorney for signing of Bid

(*Refer Clause 3.2*)

(*To be executed as an Affidavit on a Stamp paper of appropriate value*)

Know all men by these presents, We, (name of the firm and address of the registered office) do hereby irrevocably constitute, nominate, appoint and authorize Mr. / Ms (Name), son/daughter/wife of..... and presently residing at, who is presently employed with us and holding the position of, as our true and lawful attorney (hereinafter referred to as the “**Attorney**”) to do in our name and on our behalf, all such acts, deeds and things as are necessary or required in connection with or incidental to submission of our bid for the [*****] ,proposed or being developed by the [*****] (the “**Purchaser**”) including but not limited to signing and submission of all applications, bids and other documents and writings, participate in bidders' and other conferences and providing information / responses to the Purchaser, representing us in all matters before the Purchaser, signing and execution of all contracts including the rate Contract and undertakings consequent to acceptance of our bid, and generally dealing with the Purchaser in all matters in connection with or relating to or arising out of our bid for the said rate Contract and/or upon award thereof to us and/or till the entering into of the Contract with the Purchaser.

AND we hereby agree to ratify and confirm and do hereby ratify and confirm all acts, deeds and things done or caused to be done by our said Attorney pursuant to and in exercise of the powers conferred by this Power of Attorney and that all acts, deeds and things done by our said Attorney in exercise of the powers hereby conferred shall and shall always be deemed to have been done by us.

IN WITNESS WHEREOF WE,, THE ABOVE NAMED PRINCIPAL HAVE EXECUTED THIS POWER OF ATTORNEY ON THIS DAY OF, 20.....

For

(Signature, name, designation and address
of person authorized by Board Resolution
in case of Firms/Company)/Partner in case of
Partnership Firms

Witnesses:

- 1.
- 2.

Notarised
Person identified by me/personally appeared before me
/Signed before me/Attested/Authenticated*
(*Notary to specify as applicable)
(Signature, Name and Address of the Notary)
Seal of the Notary
Registration Number of the Notary
Date _____

Accepted

(Signature, name, designation and address of the Attorney)

Notes:

- *The mode of execution of the Power of Attorney should be in accordance with the procedure, if any, laid down by the applicable law and the charter documents of the executants (s) and when it is so required, the same should be under common seal affixed in accordance with the required procedure.*

Wherever required, the Bidder should submit for verification the extract of the charter documents and documents such as a board or shareholders resolution/power of attorney in favour of the person executing this Power of Attorney for the delegation of power hereunder on behalf of the Bidder.

- *For a Power of Attorney executed and issued overseas, the document will also have to be legalized by the Indian Embassy and notarized in the jurisdiction where the Power of Attorney is being issued. However, the Power of Attorney provided by Bidders from countries that have signed the Hague Legislation Convention, 1961 are not required to be legalized by the Indian Embassy if it carries a conforming Apostille certificate issued by the designated competent authority and has been notarized by the public notary.*

Annexure-8: Affidavit for blacklisting

(Non-Judicial Stamp Paper of Rs. 100/-)

Undertaking for rates, specification, blacklisting status on Stamp paper duly notarized

Tender reference No: E-248/MMGPA/ Essential Medicines

(2025-26)

1. This is to certify that the rates quoted in the bid are not higher than D.P.C.O., N.P.P.A, or not higher than MRP.
2. I/We undertake to provide the drugs/medicines/equipment's as required by Maharashtra Medical Goods Procurement Authority, Mumbai and there will be no deviation in composition, quality, packing etc.
3. The Bidder (Name of the Bidder) has not been found guilty of malpractices, misconduct or Blacklisted/Debarred/ deregistered for the quoted product by any department of Govt. of Maharashtra or by any local authority and semi Govt. organization and other State Government/Central Government's organizations/ procurement corporation as on the date of submission tender document for the quoted items.
4. I/We undertake that I/we are not involved in any major litigation such as fraud, FEMA violations and any other civil or criminal proceedings.

**Seal
Signatory**

Signature of Authorised

Date

Place

Verification

I, the above named *[Name of the Bidder]*, do hereby solemnly verify that the contents of the above Affidavit are true and correct to my knowledge and belief. Nothing false has been stated therein or material concealed therefrom.

Verified at *{location}* on *{Date}*

Note: The Bidder shall mandatorily enclose Non-Conviction Certificate issued by Licensing Authority/ State FDA along with this Affidavit for blacklisting.

Annexure- 9: Litigation Affidavit

(on Non-judicial Stamp Paper of Rs.100/-)

**Tender reference No: E-248/MMGPA/ Essential Medicines
(2025-26)**

I ----- age -----
address----- (authorized signatory to sign the
contract), hereby submit, vide this affidavit in truth, that I am the owner/authorized signatory
of the bidding entity -----and I am submitting the documents
in Envelope no.1 for the purpose of security of the contract. I hereby agree to the conditions
mentioned below:-

1. I am liable for action under Bharatiya Nyaya Sanhita (BNS) for submission of any false/
fraudulent paper/information submitted in Envelope no.1
2. I am liable for action under Bharatiya Nyaya Sanhita (BNS) if during contract period
and any false information, false bill of purchases supporting proof of purchase, proof of
testing submitted by my staff, subletting company or by myself, I will be liable for
action under Bharatiya Nyaya Sanhita (BNS).
3. I am liable for action under Bharatiya Nyaya Sanhita (BNS) if any paper is found false
/ fraudulent during contract period and even after the completion of contract
(finalization of final bill).

Authorised Signature of Bidder

Seal of Company

Verification

I, the above named [*Name of the Bidder*], do hereby solemnly verify that the contents of the
above Affidavit are true and correct to my knowledge and belief. Nothing false has been stated
therein or material concealed therefrom.

Verified at {*location*} on {*Date*}

Annexure-10: Mandate Form

01	Company Name	
02	Postal Address of the company with Telephone No., Fax No. and Mail address	
03	Name of the Managing Director/ Director/Manager Mobile No. /Phone No. E-mail address	
04	Name and designation of the authorized company official Mobile No. /Phone No. E-mail address	

Bank Details

01	Name of the Bank Branch Name & Address; Branch Code No. Branch Manager Mobile No. Branch Telephone no. Branch E-mail ID	
02	9-digit MICR code number of the bank and branch appearing on the MICR cheque issued by the bank.	
03	IFSC code of the Branch	
04	Type of Account (Current/Savings)	
05	Account Number (as appear in cheque book)	

(in lieu of the bank certificate to be obtained, please **attach the original cancelled cheque** issued by your bank for verification of the above particulars)

I/We hereby declare that the particulars given above are correct and complete. If the transaction is delayed or not effected at all for reasons of incomplete or incorrect information, I would not hold Maharashtra Medical Goods Procurement Authority, Mumbai. I have read the conditions of tender/ agreement entered and agree to discharge the responsibility expected of me/from the company as a tenderer/ successful bidder.

Date:

Company seal

Signature

Place:
designation)

(Name of the person signing &

CERTIFIED THAT THE PARTICULARS FURNISHED ABOVE BY THE COMPANY
ARE CORRECT AS PER OUR RECORDS

Bank Seal with address

Signature of the Authorized
Official of the bank

MMGPA Tender

Annexure-11: Indicative format for PRICE BID (BOQ) to be submitted online only

(To be kept in Envelope No. 2)

Item Description	Packaging	Unit	Quantity	Ex-factory cost per unit	GST applicable for Govt. Supply (In Rs.)	Other incidental charges (please specify) (In Rs.)	Total landed cost per unit (5+6+7) (In Rs.)	Total Cost Rs. (4 x8)
1	2	3	4	5	6	7	8	9

Total tender price (in words.....)

The price should be quoted only in Indian currency Note:

In case of discrepancy between unit price and total price, the unit price shall prevail. Only total landed cost per unit considered for rate comparison.

Signature of the Tenderer

Name

Designation

Business address

A separate price schedule to be used for each item while quoting rates. Each price schedule to be sealed in separate envelope mentioning PRICE BID for Item..... All such price schedule should be enclosed in envelope no. 2 which should be sealed.

To be uploaded in the form of Excel.

Annexure-12: Statement showing comparative prices offered by the tenderer in other tenders of the same product

ONLY FOR ADDITIONAL INFORMATION AS TO RATES OFFERD BY THE TENDERER IN VARIOUS OTHER TENDES.

Please mention quoted rates of above item of different years

Sr. No .	Year	MR P per unit	Price as per DPCO /NPPA	Unit Price offered in other Bids/ Tenders/Rate contracts					
				HBPC L	DMER (Govt. of Maha.)	MCG M	ESI C	Other State Govt.(s)	Tende rs of Centr al Govt.
1	2022-23								
2	2023-24								
3	2024-25								
Additional rows for information of other years can be inserted									

Signature

Seal

Schedule 1: Contract Form

(Stamp duty as applicable as per Maharashtra Stamp Act, 1958)

THIS AGREEMENT("Contract") made theday of....., 20... , at..... between Maharashtra Medical Goods Procurement Authority (MMGPA), Mumbai, formed as per the Maharashtra Medical Goods Procurement Authority Act 2023 (Mah. Act No. XIII of 2023), represented by its Chief Executive Officer and having its registered office at 1st Floor, Aarogya Bhawan, St. George's Hospital Compound, Near C.S.M.T. Railway Station, Mumbai - 400001, Maharashtra, (hereinafter "the Purchaser") of the One Part;

and

[insert name of entity], a [●] incorporated/ registered under the provisions of the [insert name of relevant statute, if applicable] and having its registered office at [●], (hereinafter referred to as the "**Supplier**" which expression shall, unless repugnant to the context or meaning thereof, include its successors, permitted assigns and substitutes) of the **OTHER PART**.

WHEREAS

- A. the Purchaser is desirous that certain specified drugs to be procured and has accepted a bid by the Supplier for the supply of [name of the drug as per the list annexed] in the sum of..... (contract price in words and figures) (hereinafter called "Contract Price").
- B. Whereas the Supplier has deposited a Bank Guarantee of Rs..... (Rs. in words.) as Performance Security towards the fulfilment of this Contract.

NOW THIS AGREEMENT 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 Supplier has accepted the Contract on the terms and condition set out in notice No.- ----- as well in the NOA Acceptance Letter No : - ----- Dt:----- which will hold good during the period of this Contract .
3. The following documents shall be deemed to form and be read and construed as part of this Agreement, viz.:
 - i. Purchase order(s) issued under this Contract, if any.
 - ii. Supplier's Acceptance to NOA
 - iii. Notification of Award (NOA)
 - iv. Supplier's Bid including response to the clarification (if any)
 - v. The Price Bid submitted by the Supplier;
 - vi. The schedule of requirements;
 - vii. The technical specifications;
 - viii. Tender Documents and all of its terms & conditions;

4. In consideration of the payments to be made by the Purchaser to the Supplier as hereinafter mentioned, the Supplier hereby covenants with the Purchaser to provide the drugs and to remedy defects therein in conformity in all respects with the provisions of the Contract.
5. The Purchaser hereby covenants to pay the Supplier in consideration of the provision of the drugs 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.
6. Upon breach by the Supplier of any of the condition of the Contract, the Purchaser may by a notice in writing, determine and terminate this Contract without prejudice to the right of the Purchaser to claim damages for antecedent breaches thereof on the part of the Supplier , as certified in writing by the Purchaser which certificate shall be conclusive evidence of the amount of such compensation payable by the Supplier to the Purchaser.
7. This Contract shall remain in force until the expiry of the date of delivery of material but notwithstanding herein or in the tender and acceptance forms contained the 'Purchaser shall not be bound to take the whole or any part of the estimated quantity herein or therein mentioned and may cancel the Contract at any time upon giving one month's notice in writing without compensating the Supplier.
8. The Supplier has fully read understood & shall abide by all the term and conditions as stipulated in Tender Document, failing which the Contract is liable to be terminated at any time without assigning any reason by the Purchaser.
9. Any change/amendments if required to be incorporated in the Contract at a later stage shall be discussed & mutually agreed by both the parties and supplementary agreements shall be binding on both the parties and shall form the part of this Contract .
10. This Contract shall be governed by and construed in accordance with the laws of Republic on India.

Brief particulars of the goods and services which shall be supplied/provided by the Supplier are as under:

Sr. No.	Brief Description Of Goods & Services	Quantity to be Supplied *	Unit Price	Total Price	Delivery Terms
					As per the supply order

***Note:**

1. Actual quantity to be supplied may vary & will be strictly as per actual requirement.
2. Actual supply to take place only after & as per the supply order(s) issued by Maharashtra Medical Goods Procurement Authority, Mumbai from time to time.
3. Tender Document is part and parcel of Contract.
4. In addition to the above, all terms & conditions as specified in Revised Manual of Office Procedures for Procurement by Government Departments: Government Resolution no.

Bha.kha.sa-2014/ Pra. Kra. 82/Section-III/Industry-4, dated 1.12.2016 issued by Department of Industries and Labour, Government of Maharashtra will apply on the Contract.

IN WITNESS whereof the parties hereto have caused this Contract to be executed in accordance with their respective laws the day and year first above written.

Signed, Sealed and Delivered
(on behalf of the Purchaser)

Signed, Sealed and Delivered
(on behalf of the Supplier)

Address for communication:

Address for communication:

In presence of

1.

2.

Office of the-

Sd/-

**Chief Executive Officer,
Maharashtra Medical Goods
Procurement Authority, Mumbai**

**1st Floor, Aarogya Bhawan, Near
CSMT Railway Station, Mumbai
400001 (Maharashtra)**

Schedule-2 Performance Security Form

To,

**Chief Executive Officer,
Maharashtra Medical Goods Procurement
Authority,
1st Floor, Aarogya Bhawan,
Near CSMT Railway Station,
Mumbai 400001 (Maharashtra)**

Dear Sir

Whereas you intend to enter into a contract, as per your Notification of Award, Reference No. _____ dated _____ (hereinafter referred to as "the contract") with M/s _____ as Supplier for the supply of _____ defined in contracts schedule, (hereinafter referred to as "drugs") and whereas the Supplier has undertaken to produce a performance cum warranty bond for amount of Rs _____ being equal to 3% of the total contract value of the drugs to be delivered as specified in NOA No _____ dated _____.

1. We _____ (Name of the Bank), hereby expressly, irrevocably, and unreservedly undertake and guarantee as principal obligators on behalf of the Supplier that in the event that the Maharashtra Medical Goods Procurement Authority ("MMGPA") submits a written demand to us stating that the Supplier has not performed according to the terms and conditions of the contract, we will pay MMGPA on demand and without demur any sum up to a maximum amount of (3% of the contract value). Any claims must bear the confirmation of MMGPA's bankers that the signatures thereon are authentic. MMGPA's written demand shall be conclusive evidence for us to make payment to MMGPA. For the avoidance of doubt, any documents received by way of facsimile or similar electronic means is/are not acceptable for any purpose(s) under this guarantee.
2. We shall not be discharged or released from this undertaking and guarantee by any arrangements, variations made between beneficiary and the seller or any forbearance whether as to payment, time performance or otherwise.
3. Unless a demand under this guarantee is received by us in writing on or before the expiry dates (unless this guarantee is extended by the Supplier), all MMGPA's rights under this guarantee shall be forfeited and we shall be discharged from the liabilities hereunder.
4. This guarantee shall be a continuing guarantee (which means guarantee will also be valid if the bank is in under liquidation or bankruptcy) and shall not be discharged by any change in the constitution of the bank or in the constitution of the Supplier.
5. Please return this letter of guarantee immediately after our liability thereafter has ceased to be valid.
6. Our liability under this guarantee will cease to be valid even if the guaranteed deed is not returned to us.
7. This guarantee is personal to MMGPA and not assignable to a third party without our prior written consent.

8. This guarantee shall be governed by Indian Law. This guarantee is valid until <<mention date { date of validity should not be less than 24 months from signing of the Contract} >>.

Signature and Seal of Guarantors _____

Date _____

Address: _____

UNREPAIRED

Schedule-3: Supply Schedule

(in accordance with clause 4.10)

Dt:

Supplier Name:

PO No:

PO Date:

Drug Name:

Total Ordered

Qty:

Sr.No.	Consignee Name	Ordered Qty	Expected Date of drug to be manufactured at manufacturing unit	Expected Date of in house/NABL testing	Expected Date of Delivery	No. of days taken from PO Date	Remarks, if any
1.							
2.							
3.							

Seal

Signature

Date
Place

Schedule 4: Schedule for Packing of Drugs and Medicines

A. General Specifications: All drugs should be packed & supplied in prescribed packing only.

1. No corrugate package should weigh more than 15 Kgs (i.e., Product + Inner Carton Corrugated box)
2. All corrugated boxes should be of 'A' grade paper .i.e. Virgin.
3. All items should be packed only in first hand boxes only.
4. Flute - The corrugated boxes should be of narrow flute.
5. Joint - Every box should be preferably single joint and not more than two joints.
6. Stitching - Every box should be stitched using pairs of metal pins with an interval of two inches between each pair. The boxes should be stitched and not joined using calico at the corners.
7. Flap - The flaps should uniformly meet but should not overlap each other. The flap when turned by 45 - 60 degree should not crack.
8. Tape - Every box should be sealed with gum tape running along the top and lower opening.
9. Carry strap- Every box should be strapped with two parallel nylon carry straps (they should intersect).
10. Label - Every corrugated box should carry a large outer label clearly indicating that the drug is for '**Government of Maharashtra (MMGPA) Supply - Not for Sale**', wherein '**Government of Maharashtra (MMGPA) Supply**' should be in readable purple colour and '**Not for Sale**' should be in green colour
11. The product label on the carton should be large at least 15 cm. x 10 cm. dimension. It should carry the correct technical name, strength or the drug, date of manufacture & distributor, date of expiry, quantity packed and net weight of the box.
12. Other - No box should contain mixed drugs or mixed batches of the same drug.

B. Specification for Corrugated Boxes Holding Tablets / Capsules / Pessaries

1. The box should not weigh more than 7-8 Kilograms. The grammage of outer box should be 150 gsm and inside partition / lining should be 120gsm.
2. The box should be of 5 ply with bursting strength of 9 Kg/ Cm²

C. Specifications for Ointment / Cream / Gels Packed in Tubes

1. No corrugate box should weigh more than 7-8 Kgs.
2. Every Ointment tube should be individually packed in carton and then packed in 20's in a White board box, which may be packed in a corrugated box.
3. The grammage of outer box should be 150 gsm and inside partition / lining should be 120gsm.

D. Specifications for Injectable (in Vials and Ampoules)

1. Vials may be packed in corrugated boxes weighing up to 15 Kilograms. Ampoules should be packed in C.B weighing not more than 8 Kilograms.
2. Corrugated box for vials should be of 150 gsm (outer box should be 150 gsm and inside partition / lining should be 120 gsm) and 7 ply, while corrugated box for

ampoules should be of 150 gsm (outer box should be 150 gsm and inside partition / lining should be 120 gsm) and 5 ply.

3. Bursting strength for CB boxes for
 - a. Vials : Note less than 13 Kg/Cm²
 - b. Amp : Note less than 9 Kg/Cm²
4. In the case of 10 ml Ampoules, 100 or 50 ampoules may be packed in a White board box. Multiples of White board boxes packed in corrugated box. In case of Ampoules larger than 10 ml, only 25 ampoules may be packed in a White board box with partition.
5. If the vial is packed in individual carton, there is no necessity for White board box packing. The individual carton may be packed as such in the corrugated box with centre pad.
6. In case of ampoules, every White board box should carry 5 amps. Cutters placed in a polythene bag.
7. Vials of Eye and Ear drops should be packed in an individual carton with a dispensing device. If the vial is of FFS/BFS technology, they should be packed in 50's in a White board box.

E. Primary Package

1. Tablets & Capsules

- 10 Tablets/Capsules or multiples of 10 should be packed in an Aluminum strip / Aluminum – P V C blister pack.
- Aluminum strips: Thickness of Aluminum foil: 40 microns with LDPE 25-micron coating/heat seal lacquer
- PVC Film : Transparent, clear/amber, food grade, blister forming PVC film, Film gauge 200 microns, P E coating : 25 microns, PVdC coating : 60 gsm.
- Aluminum foil: Hard tempered Blister foil, VMCH coated, Thickness: 0.025 mm.

'Government of Maharashtra (MMGPA) Supply - Not for Sale' in readable purple or green colour on each strip.

2. Injections

- Injection in ampoule form should be supplied in Double constructed neck ampoules with the label bearing the words **'Government of Maharashtra (MMGPA) Supply - Not for Sale' in readable purple or Green colour.**
- The vials should be supplied with Aluminum seal ampoules with the label bearing the words **'Government of Maharashtra (MMGPA) Supply - Not for Sale' in readable purple or Green colour.**

3. Liquid Orals

Liquid preparations should be in FDA approved glass/plastic bottles with pilfer-proof caps. The top of the cap and the label to be affixed on the containers should bear a distinct colour different from the colour of the label of the trade packs and they should be overprinted in readable purple colour with the words **'Government of Maharashtra (MMGPA) Supply - Not for Sale'.**

4. Ointments

Ointments should be supplied in tube and bearing the words '**Government of Maharashtra (MMGPA) Supply - Not for Sale**' in readable purple or green colour.

F. Secondary Package

The strips/ampoules/vials, tubes and bottles should be packed in boxes for easy handling, transport and distribution. It shall be fabricated from Millboard/grey board/cardboard with appropriate bursting strength. The secondary packaging material must be clearly labelled with the names of item, batch number, mfg date, expiry date and the number of units per box. The secondary box shall bear the words '**Government of Maharashtra (MMGPA) Supply - Not for Sale**' in readable purple or Green colour.

G. Tertiary Package

The boxes shall be packed in weather resistant triple walled insulated corrugated 5 ply cartons, each ply having strength of minimum 150 gsm It should be fabricated from virgin quality "A" grade material. The overall dimension of the carton should be such that the drug does not get damaged during transportation and storage. Each international shipping carton should weigh less than 50 kg. It is important that individual boxes are not too heavy during transport as they are frequently loaded and off loaded manually at airports and intermediate stores.

H. Case Identification

All cases should prominently indicate the following:

1. Purchaser's line and code numbers
2. The generic name of the drug
3. The dosage form (Tablet, Ampoule, Syrup)
4. Date of manufacture and expiry (month and year) (in clear language not code)
5. Batch number
6. Quantity per case (Carton containing----- secondary packages)
7. Special instructions for storage and handling
8. Name and address of manufacture
9. Any additional cautionary statements.

SPECIMEN LABEL FOR OUTER CARTON (20cm x 15cm)

Government of Maharashtra (MMGPA) Supply - Not for Sale

~~~~~

Generic Name of Drug I.P.

~~~~~

10 x 10 TABLETS

Batch No: XXXXXX

Quantity Packed: XXXXXX

Mfg. Date: XXXX- 2025

Exp. Date: XXXX -2027

Manufactured by: M/s. XXXXXX

Carton containing -----secondary packages

Special instructions for storage and handling - Store in a Cool and Dry Place

Bar Code

WINGPATENTED

Schedule 5 Prebid Queries Format

Name of the Bidder :

(A): Prebid Query Format for Tender Terms & Conditions

Sr no	Tender Page no	Tender Clause no	Clause Title	Query / Clarification	Justification by Bidder
1					
2					
3					
4					

(B): Prebid Query Format for Itemwise (Medicines)

Sr no	Item Sr no (As per Appendix A)	Name of the Item	Specifications	Query / Clarification	Justification by Bidder
1					
2					
3					
4					