

MAHARASHTRA MEDICAL GOODS PROCUREMENT AUTHORITY

Expression of Interest (EOI) for Market Assessment of Original Manufacturers, Authorized Importers of Equipments (Year 2026-27)

**EOI reference No: E- 297/MMGPA/ EOI for Equipments
(2026-27)**

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Near C.S.M.T. Railway Station, Mumbai - 400 001.
Maharashtra

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Disclaimer

This Expression of Interest (EOI) Document is not an agreement and is neither an offer nor invitation by the MMGPA to the prospective EOI Responders or any other person. The purpose of this Expression of Interest (EOI) Document is to invite interested and eligible entities to provide information relating to their regulatory status, manufacturing capability, quality systems, market presence, and related parameters, solely to enable MMGPA to undertake a market assessment and planning exercise in connection with any potential future procurement of equipment.

This Expression of Interest (EOI) Document contains information, statements, and indicative details prepared by MMGPA solely for the purpose of facilitating a broad market assessment and planning exercise in relation to the potential future procurement of equipment. Such information and statements are based on assumptions, preliminary assessments, and internal considerations and are not intended to be exhaustive or definitive. This EOI Document may not address the specific circumstances, operational capacities, or regulatory positions of every interested entity, and MMGPA, its employees, or advisors do not undertake to consider the individual requirements of each person or entity reviewing or responding to this document. The information provided herein may be subject to change and may not be complete, accurate, or current in all respects. Each EOI Respondent is therefore advised to conduct its own assessment, verification, and analysis of the information contained in this EOI Document and to seek independent professional or technical advice as it considers appropriate.

Information provided in this Expression of Interest (EOI) Document to the EOI Responder(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 EOI Responder(s) 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 Expression of Interest (EOI) Document or otherwise, including the accuracy, adequacy, correctness, completeness or reliability of the Expression of Interest (EOI) Document and any assessment, assumption, statement or information contained therein or deemed to form part of this Expression of Interest (EOI) Document or arising in any way for participation in this Expression of Interest (EOI) 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 Expression of Interest (EOI) Document.

The EOI Responder(s) shall bear all its costs associated with or relating to the preparation and submission of its EOI Response 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 EOI Response. All such costs and expenses will remain with the EOI Responder(s) 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 EOI Responder(s) in preparation or submission of the EOI Response, regardless of the conduct or outcome of the responding process. The MMGPA also accepts no liability of any nature whether resulting from negligence or otherwise howsoever caused arising from reliance of

any EOI Responder(s) upon the statements contained in this Expression of Interest (EOI) Document. MMGPA shall not be responsible or liable for any costs, expenses, or losses incurred by EOI Respondents in connection with preparation or submission of EOI Responses, regardless of the outcome of the EOI process. Participation in this Expression of Interest (EOI) shall not confer any right or entitlement on the EOI Respondents to participate in any subsequent procurement process.

MMGPA reserves the right to independently determine the eligibility criteria, qualification requirements, and procurement strategy in any future tender process, without any reference to or obligation arising from this EOI.

Abbreviations and Glossary

Abbreviations	Description
BG	Bank Guarantee
BOM/BOQ	Bill Of Material/Quantity
CA	CHARTERED ACCOUNTANT
CAMC	Comprehensive Annual Maintenance Contract
CBS	Cost Based Selection
CMC	Comprehensive Maintenance Contract
CRC	Consignee Receipt certificate
DPIIT	Department for Promotion of Industry and Internal Trade
EMD	Earnest Money Deposit
EM-II	Entrepreneurs Memorandum
FEMA	Foreign Exchange Management Act
GST	Goods and Services Tax
IA	Implementation Agency
IP	Intellectual Property
IQ	Installation Qualification,
ISO	International Organization of Standardization
KPI	Key Performing Indicators
LLP	Limited Liability Partnership
MMGPA	Maharashtra Medical Goods Procurement Authority
MSME	Ministry of Micro, Small & Medium Enterprises
NEFT	National Electronic Funds Transfer
O&M	Operation and Maintenance
OEM	Original Equipment Manufacturer
OP	Operational Qualification
PAN	Permanent Account Number
PO	Purchase Order
PQ	Performance Qualification
RFP	Request For Proposal
RTGS	Real Time Gross Settlement
SSI	Small-scale industries
TCV	Total Contract Value

Notice and Schedule Inviting Expression of Interest (EOI)

EOI Notice

The Maharashtra Medical Goods Procurement Authority (MMGPA) invites Expressions of Interest (EOI) from eligible Original Manufacturers(OEM's), Authorized Importers for assessing market capability, regulatory compliance, manufacturing capacity, and delivery capability for supply of equipment to public health institutions in the State of Maharashtra.

This EOI is a non-binding, pre-procurement exercise and does not constitute a tender or offer of contract. No price submissions, financial offers, or securities are required at this stage.

The objective of this EOI is to assess the number of eligible and capable OEMs, Authorized Importers available for the listed essential items and their readiness to participate in future procurement processes.

The information collected through this EOI will be used for planning future procurement strategies. MMGPA reserves the right to initiate or not initiate any subsequent tender or rate contract process without any obligation to the EOI respondents.

Only Original Manufacturers or their Authorized Importers are eligible to participate.

Interested entities meeting the eligibility conditions are invited to submit their EOI responses within the timelines specified in this document.

EOI responses must be submitted within fifteen (15) days from the date of publication of this EOI. Late submissions shall not be accepted.

All response-related activities, including downloading of documents and submission of responses, shall be governed by the schedule indicated below.

EOI Response Schedule

No.	Activity	Date & Time
1	Publication of Notice Inviting Expression of Interest (EOI)	From 01.06.2026 at 11:00 AM
2	Last date and time for submission of EOI Responses	15.06.2026 at 2:00 PM
3	Opening of EOI Responses (Technical & capability only, No price information shall be submitted)	dt.16.06.2026 at 3:00 PM
4	Eoi Clarification Meeting	From 08.06.2026 , 02:00 PM
5	Address for communication	1st Floor, Aarogya Bhawan, St. Georges Hospital Compound, Near CSMT Railway Station, Mumbai- 400 001

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

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 authority 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 as amended from time to time.

- 1.1 All related activities (process) like EOI downloading and response submission will be governed by the EOI Schedule.
- 1.2 All activities of this EOI are carried out online on website <https://mahatenders.gov.in>. The EOI 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.
- 1.3 No commitment is made by MMGPA regarding procurement of any quantity pursuant to this EOI, and the Authority reserves the right to modify, expand, curtail, or altogether omit any items or quantities in any subsequent procurement process, if undertaken
- 1.4 EOI 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

Section 2: General Definitions

For the purposes of this Expression of Interest (EOI) Document, and unless the context otherwise requires, the following terms shall have the meanings assigned to them below:

- 2.1 **“Applicable Laws”** Means all laws, statutes, rules, regulations, notifications, guidelines, and policies in force in India as issued by the Government of India or the Government of Maharashtra, including but not limited to the Drugs and Cosmetics Act, 1940 and Rules thereunder, Medical Device Rules, 2017, New Drugs and Clinical Trials Rules, 2019, NPPA/DPCO provisions, and other applicable regulatory requirements, as amended from time to time.
- 2.2 **“Authority” or “MMGPA”** Means the Maharashtra Medical Goods Procurement Authority, established under the Maharashtra Medical Goods Procurement Authority Act, 2023.
- 2.3 **“Expression of Interest” or “EOI”** Means this non-binding, pre-procurement process initiated by MMGPA to solicit information regarding market capability, regulatory compliance, manufacturing capacity, and quality systems from eligible entities.
- 2.4 **“EOI Document”** Means this document, including all sections, appendices, annexures (if any), and amendments issued by MMGPA in connection with the Expression of Interest.
- 2.5 **“EOI Response”** Means the information and declarations submitted by an EOI Respondent in response to this EOI Document, limited to technical, regulatory, manufacturing, and capability-related information.
- 2.6 **“EOI Respondent”** Means any manufacturer or importer who submits an EOI Response pursuant to this EOI Document.
- 2.7 **“Conflict of Interest”** Means any situation in which an EOI Respondent has interests that could improperly influence its participation in the EOI process, including but not limited to relationships that compromise fairness, transparency, or objectivity.
- 2.8 **“Fraudulent or Corrupt Practices”** Means misrepresentation, concealment of facts, submission of false or forged documents, or any attempt to influence the EOI process improperly.
- 2.9 **“Government”** Means the Government of India or the Government of Maharashtra, as applicable.

Interpretation Words and expressions used but not defined in this EOI Document shall have the meanings assigned to them under Applicable Laws.

Note: The words and expressions used in this EOI, 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 the EOI and contains the following terms & conditions governing the EOI.

3.1 Responsibility of verification of contents of EOI

- 3.1.1 It shall be the responsibility of the EOI Responders to read/examine all instructions, forms, terms, conditions, specifications, schedules, appendix, annexures and other provisions contained in the Expression of Interest (EOI) Document, including any addendum or corrigendum issued thereunder prior to submission of the response.
- 3.1.2 Responders shall ensure that all required documents as specified in **Appendix A (Checklist)** are duly uploaded.

3.2 Acknowledgement of the Responder

- 3.2.1 It shall be deemed that by submitting the EOI, the Responder has:
- i. examined the EOI Document and understood its contents.
 - ii. received all relevant information available from MMGPA
 - iii. acknowledged and agreed that inadequacy, lack of completeness or incorrectness of information provided in the Expression of Interest (EOI) Documents shall not be a basis for any claim for compensation, damages or extension of time.
 - iv. acknowledged that it does not have a Conflict of Interest as per Clause 3.9 and
 - v. agreed to be bound by the provisions and undertakings provided by it under and in terms hereof.

MMGPA 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 EOI or its Responding process, including any error or mistake therein or in any information or data given by MMGPA.

3.3 Period of Validity of EOI

- 3.3.1 The EOI Response submitted pursuant to this document shall remain valid for the purpose of this EOI process only until completion of the EOI exercise or initiation of any subsequent procurement process, whichever is earlier.

3.4 Nature of the Expression of Interest

- 3.4.1 The information received pursuant to this EOI shall be used by MMGPA only for:
- a) market assessment and planning,
 - b) identification of technically and regulatorily suitable entities, and
 - c) formulation of future procurement strategies, if any.
- 3.4.2 This Expression of Interest (EOI) is a non-binding, pre-procurement exercise issued solely for the purpose of assessing market capability, regulatory compliance, manufacturing capacity, and quality systems of eligible entities.

- 3.4.3 Submission of an EOI Response does not constitute a bid, tender, or offer, and does not create any contractual, financial, or legal obligation on either MMGPA or the EOI Respondent.
- 3.4.4 MMGPA reserves the right to initiate or not initiate any subsequent procurement process, including a tender or RFP, without assigning any reason.
- 3.4.5 The EOI is intended to assess the number of eligible and capable Original Manufacturers and Authorized Importers available for each essential equipment/item and their readiness to participate in future procurement processes.

3.5 Responsibility of EOI Respondents

- 3.5.1 Each EOI Respondent shall:
- a) carefully examine the EOI Document and all instructions therein;
 - b) ensure that all information and documents submitted are true, accurate, and complete to the best of its knowledge;
 - c) satisfy itself regarding the scope, intent, and implications of the EOI.
 - d) Any misrepresentation, concealment, or submission of false information may result in rejection of the EOI Response.
- 3.5.2 Respondents shall provide accurate information regarding:
- (a) availability of equipment/item(s) against the provided list,
 - (b) manufacturing/import capacity,
 - (c) indicative delivery timelines.

3.6 Submission of EOI Responses

- 3.6.1 EOI Responses must be submitted within fifteen (15) days from the date of publication of the EOI. Late submissions shall not be accepted.
- 3.6.2 The EOI Response shall be limited to technical, regulatory, manufacturing, and capability-related information only.
- 3.6.3 No price bids, financial offers, bid securities, earnest money deposits, or payments of any kind are required or permitted at the EOI stage.

3.7 Clarifications and Amendments

- 3.7.1 EOI Respondents may seek clarification, within the timelines specified in the EOI Schedule.
- 3.7.2 MMGPA may, at its discretion:
- a. issue clarifications,
 - b. amend the EOI Document, or
 - c. modify the EOI Schedule.
- 3.7.3 All such clarifications or amendments shall form an integral part of the EOI Document.

3.8 Conflict of Interest

- 3.8.1 EOI Respondents shall disclose any actual or potential conflict of interest that may affect their participation in the EOI process.
- 3.8.2 MMGPA reserves the right to reject an EOI Response where a conflict of interest is found to compromise fairness, transparency, or objectivity.

3.9 Fraudulent and Corrupt Practices

- 3.9.1 EOI Respondents shall observe the highest ethical standards during the EOI process.
- 3.9.2 Any attempt to:
 - a. misrepresent facts,
 - b. submit forged or misleading documents, or
 - c. improperly influence the EOI process
- 3.9.3 may result in rejection of the EOI Response and may render the entity ineligible for participation in future EOIs issued by MMGPA.

3.10 Confidentiality

- 3.10.1 Information submitted by EOI Respondents shall be treated as confidential to the extent permitted under applicable laws.
- 3.10.2 MMGPA may disclose such information where required by law, regulation, or directions of statutory or judicial authorities.

3.11 Governing Law

This EOI process shall be governed by and construed in accordance with the laws of India.

3.12 Manufacturing/Importing and Product Permissions

- 3.12.1 EOI Respondents shall furnish details of their regulatory status as pharmaceutical manufacturers and/or importers licensed under the provisions of the Medical Device Rules 2017, Drugs and Cosmetics Act, 1940 and the rules made thereunder. Such details may include manufacturing licenses issued by the competent State Licensing Authority (SLA) and/or Central Licensing Authority (CLA), including licenses in Forms 25, 28, 28D, 28E, MD-5, and MD-9, as applicable to the drugs proposed, together with validity or retention particulars.
- 3.12.2 In the case of importers, Respondents shall furnish details of import licenses issued by the Drugs Controller General of India (DCGI), including Forms 10 or 10-A, along with relevant import registrations such as Form 41 or Form MD-15, and associated wholesale licenses such as Forms 20B and/or 21B, wherever applicable.
- 3.12.3 In respect of New Drugs and/or Fixed Dose Combinations, Respondents shall furnish details of import and/or manufacturing permissions obtained under the Drugs and Cosmetics Act, 1940 and the New Drugs and Clinical Trials Rules, 2019, including permissions such as Forms CT-20 and/or CT-23, where applicable.

3.12.4 In case of imported products, one Original Manufacturer can authorized one Agent.

3.12.5 Respondents shall also indicate linkage between manufacturing/import approvals and supply capability, including delivery timelines.

3.13 Delivery Capability

3.13.1 EOI Respondents shall provide indicative details on:

- (a) production lead time,
- (b) dispatch timeline,
- (c) emergency supply capability,
- (d) monthly production capacity,
- (e) warehousing/logistics arrangements in Maharashtra.

3.14 Respondent Classification

3.14.1 EOI Respondents shall provide information on:

- (a) MSME status,
- (b) SC/ST ownership,
- (c) location – within Maharashtra / outside Maharashtra.

3.15 Eligibility Criteria

Sr. No.	Basic Requirement	Specific Requirement	Documents required
1.	Registered Legal Entity	The Respondent shall be any person/Company/ Society/Proprietorship/ Partnership firm/Trust registered under applicable Act in India/ Government-owned enterprise or institution The Respondent shall be – a) A manufacturer having valid manufacturing and equipment license for the items quoted. OR b) An Importer* having valid import license and equipment license for the items quoted. c) Registered with the GST Authorities. d) Should have a valid PAN & TAN number. Proprietorship Firms are not allowed to participate in this EOI. <i>*Importer refers to a legal Entity such as a Company/ Society/ Trust/Partnership firm registered</i>	a. 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. b. Attested photocopy of valid manufacturing Equipment/ import license with product list duly approved by the Licensing 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. If quoted item is manufactured at different

Sr. No.	Basic Requirement	Specific Requirement	Documents required
		<i>under applicable Act in India/ Government-owned enterprise or institution that engages in the process of bringing equipment or goods from outside India into the country's borders for commercial purposes. Importer itself shall be responsible for supply and maintenance of the equipment as per the terms of RFP and shall not engage any third party for the same)</i>	places, Manufacturing License & Performance certificate should be enclosed. c. Manufacturer's Authorization as per Annexure XIV to be provided by Importer d. Copy of GST Registration certificate issued by GSTN authorities. In case of e. Copy of PAN Card.
2.	Certifications/ registration	The Respondent shall have to provide requisite certifications/registration.	a. Certificates of DPIIT (if applicable) b. Original manufacturer's certificate that the product is being used in country of origin. c. Import Export Certificate (IEC Code) d. Affidavit of Importer regarding equipment being imported in India for last three years.
3.	Litigation	The Respondent 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 VII.
4.	Conflict of Interest	On the date of submission of the proposal, the Respondent should not be involved in any conflict-of-interest situation.	Undertaking by the authorized signatory as per Annexure I
5.	Blacklisting or banned	On the date of submission of the proposal, the Respondent should not be blacklisted or banned 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.	Affidavit as per Annexure VII
6.	Debarment	On the date of submission of the proposal, the Respondent should not	Affidavit as per Annexure VII.

Sr. No.	Basic Requirement	Specific Requirement	Documents required
		be debarred	
7.	Average Annual Turnover	The Average Annual Turnover of the Respondent for the last three financial years (2022–23, 2023–24, 2024–25) shall be equal to or greater than the minimum turnover requirement specified in Schedule of Requirement (Appendix A) for each equipment/item for which the Respondent submits an EOI.	Certificate issued by a statutory auditor/chartered accountant (as attached Annexure-IV) along with Audited Financial Statements confirming the Average Annual Turnover of the Respondent during the stated Financial Years must be submitted.
8.	Net Worth	The net worth of the Respondent in the financial year immediately preceding the EOI Due date should be positive and should have not eroded by more than 30% (thirty percent) in the last three (3) years.	Certificate issued by a statutory auditor/chartered accountant (as attached Annexure-IV).
9.	Technical Capability	The Respondent must have successfully completed supply, installation, and commissioning of the same or similar equipment during the last three financial years (2022–23, 2023–24, 2024–25), with a cumulative quantity equivalent to at least eighty percent (80%) of the quantity specified in Appendix A for the respective equipment/item(s) for which the Respondent submits an EOI. The Respondent shall submit the work done with satisfaction from the concerned authority. For the purpose of quantification of 80% similar work, only the Purchase order from Govt./Semi Govt.& Hospitals registered under Maharashtra Public Trust Act 1950 shall be considered. Purchases order from private hospitals shall not be considered.	The Respondent shall provide the documentary evidence in support of its credentials such as agreement copy/ work order / Letter of Award. This should be supported by work completion certificate/customer satisfaction certificates with customer details and client satisfactory certificate. Statutory auditor's certificate or Chartered Accountant's certificate, as the case may be, shall be submitted for demonstrating the past experience. (as per Annexure number III)

Sr. No.	Basic Requirement	Specific Requirement	Documents required
10.	Production Capacity / Import Quantity	For Original Equipment Manufacturers (OEMs), the Production Capacity in the last financial year for the relevant equipment/item(s) shall be at least two (2) times the quantity specified in Appendix A. In case of Importers, the cumulative quantity imported/supplied in the last financial year for the relevant equipment/item(s), or the production capacity of the OEM (supported by certification), shall be at least two (2) times the quantity specified in Appendix A.	Certificate of Statutory Auditor/Chartered Accountant For importers ,Certificate of Statutory Auditor/Chartered Accountant of OEM has to be submitted in Annexure III
11.	Turn around time for service of equipment	The respondent shall attend all service/break down complaints within 24hrs.of complaint registration. The reported fault shall be rectified and the equipment restored to normal working condition within 48hrs of complaint registration.	-
12.	Number of service station	The respondent shall have sufficient number of service centres so that the non functional equipments should be rectified with 48 hrs.from the receipt of complaint	Proof of locations of service stations must be submitted as a part of EOI
13.	Proof of Distribution Network	Respondents, 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 EOI submission. In case there is no registered office or established operational distributional network the respondent shall give undertaking for establishment of both within three (3) months from the last date of submission of EOI.	Proof of distribution network— such as a valid drug/product/item(s) selling/distribution license issued by the state's licensing authority, along with details (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
14.	Quantity specified in EOI	The quantity mentioned in the EOI is indicative in nature. Detail requirement will be given in subsequent RFP.	-

Appendix A: Schedule of Requirements

Appendix A-Schedule of requirement			
SR.NO	EQUIPMENT NAME	Estimated Quantity	Annual Turnover
1	Multipara Monitor	3566	30,38,44,500
2	ICU Ventilator	392	41,18,58,000
3	Biphasic Detibrillator	131	5,58,54,000
4	Central Monitor System for 20 BED	151	45,94,86,000
5	Multipara Monitor	1370	11,67,45,300
6	central Monitor System for 20 BED ICU (1x20 Multipara Monitor)	27	8,22,96,000
7	Patient stretcher (fully SS)	326	49,32,900
8	Wheel Chair Foldable	326	26,03,475
9	Crash cart	261	1,35,92,880
10	Food step-Double	3566	1,19,82,600
11	Beside locker	3566	2,09,69,550
12	IV stand with SS Rod and Castor Base	3566	86,87,385
13	Biomedical Waste Small set of 3	3566	89,86,950
14	Dressing Trolley	261	61,38,720
15	Dressing Trolley	261	39,46,320
16	ECG Machine Trolley	216	32,65,920

17	Instrument Trolley	216	34,47,360
18	Oxygen Cylinder Trolley	261	8,55,036
19	Infusion Pump	3566	17,97,39,000
20	ECG Machine 12 Channel	196	1,56,20,850
21	ABG Machine with ISE ion	131	15,24,50,100
22	Syrings Pump	3566	14,37,91,200
23	Bed side Stool	3566	1,10,83,905
24	ICU Bed Motorized With Mattress	3566	32,95,21,500
25	Overbed Table	3566	3,29,52,150
26	Intusion Pump	205	1,03,19,400
27	Syringe Pump	1665	6,71,32,800
28	ABG Machine with ISE	65	7,62,25,050
29	Chesa INC	329	11,51,06,400
30	Iotome Electric India Pvt.Ltd.	736	19,30,60,800
31	X-ray Machine 500 MA	104	18,20,56,500
32	X-ray Machine 300 MA	128	12,20,42,700
33	X-ray Machine 100 MA	104	6,61,28,220
34	Sonography Machine	304	63,14,48,379
35	Ultraportable Handheld Digital X ray	180	88,12,80,000

36	Phaco Machine	135	35,91,00,000
37	ECG Machine	279	3,31,22,880
38	Sonography M/c	7	1,60,86,600
39	Neonatal Pulse Oximeter	376	1,49,51,844
40	Handheld Pulse Oximeter	3985	4,59,04,320
41	Fetal Monitor/NST Machine	740	11,81,14,290
42	PhotoTherapy	203	80,59,500
43	Radiant Warmer	1231	9,79,18,470
44	Fetal Doppler	5463	2,11,96,440
45	CPAP Machine	63	2,00,83,644
46	Neonatal Monitor	16	1,76,20,470
47	T Piece resusciator	378	4,51,21,709
48	Syrings Pump	297	93,25,800
49	Neonatal Monitor	92	98,85,510
50	Multipara Monitor(ADULT)	279	3,54,77,640
51	LED Slit lamp	63	1,92,27,600
52	Surgical set for contract	2585	5,63,56,382
53	Auto Refract k.	122	3,70,81,800
54	A Scan Biometer	45	78,47,964

55	Portable Funds Camera	56	1,71,67,950
56	Non. Contact Tonometer	68	2,35,43,946
57	Operating Microscope	56	7,35,74,955
58	Rectam for Pediatric Screeing	11	2,45,24,991
59	Mobile X-Ray	11	1,25,92,843
60	Syringe Pump	281	71,55,540
61	Monitor	56	69,16,493
62	Pulse Oximeter	23	9,25,384
63	CPAP	68	2,48,70,767
64	Pulse oximeter adult padiatric and neonatal probes	281	80,84,700
65	Power Bone Drill	32	2,64,41,100
66	Pneumatic Tourniquet	25	81,92,923
67	Cardiac Cath Lab with Tumkey	5	39,06,00,000
68	Handheld Pulse Oxymeter	14931	18,22,77,648
69	Laproscope	254	22,17,05,797
70	Non Invasive Ventilator	16	1,23,60,587
71	Pediatric Dental Chair	72	91,41,350
72	Wall Mounted dental X-Ray	72	1,25,56,742
73	Digital Radiography with Tumkey	2	2,34,00,000
74	Anesthesia Workstation	5	1,17,72,000
75	Ventilator (Adult+Peed)	2	41,40,000
76	Ventilator (Neonate)	2	41,40,000
77	UV -Visible Spectrophotometer with attached CPU and Printer	79	5,86,37,313
78	ION Selective Meter (iono meter) wiith specific electrode	68	2,35,44,054
79	Quartz Double ditillation Unit	34	63,08,280
80	Soxhelet Extraction unit (Closed	2	11,06,278

	System)		
81	Kjeldahl digestion and distillation unit	18	1,26,25,906
82	Biosafety Cabinet BS-2	45	1,76,58,000
83	Digital Weighing Balance	45	58,64,400
84	GC with auto sampler +detector ECD,FPD,FID	7	2,37,16,584
85	AAS+Hybrid Generator+19HC lamps+accessories+Consumables	7	2,07,14,098
86	Microwave Digester8-12 tubes	7	1,48,09,495
87	Double dome shadowless ceiling mounted OT lights	77	3,31,38,698
88	Shadow OT lights mobile	227	2,93,95,242
89	C-Arm Compatible Operation table	34	3,51,24,840
90	OT Table ordinary	45	38,66,940
91	Brain Sanner Machine	77	10,40,40,000
92	PO Retinal Camera	52	11,28,14,959
93	Power Bone Drill	23	1,88,86,500
94	Pneumatic Tourniquet	38	1,26,61,790
95	C-Arm Compatible Operation table	29	3,04,41,528
96	Fetal Monitor/Non Stress Test Machine	257	3,35,49,995
97	Autoclave HP Horizontal	54	2,35,43,957
98	Trolley Stretcher	569	59,56,177
99	Wheel chairs	326	51,20,820
100	Cell Counter	2500	68,98,31,010
101	Microscope	2687	16,31,24,280
102	Lab Autoclave	1852	2,19,89,902
103	Hemoglobinostrips Compatible with Hemoglobinometer	1400119	2,00,72,102
104	Digital Hemoglobinostrips Compatible with Hemoglobinometer	7331	3,61,24,704

105	Examination Table with Lithotomy Position Arrangement	2529	5,00,43,852
106	Lamp with Stand	2061	2,67,93,000
107	Binocular Microscope	288	56,70,144
108	LED FM Microscope	68	38,01,600
109	Objective lense 100x (oil immersion	308	5,91,840
110	Microscope eye piece 10x	457	5,11,560
111	OAE Surenet Machine	1071	34,22,05,920
112	Bera with ASSR	45	3,78,72,000
113	Sterile connecting device	52	5,32,50,750
114	MGIT	2	1,12,24,147
115	Twincubator	7	56,70,000
116	GTBLOT 48	11	2,16,00,000
117	Glucometer	3137	16,30,980
118	ION Selective Electrode Electrolyte Analyzer	3071	1,14,18,90,750
119	ESR Analyzer Availabilty	3290	1,33,52,73,840
120	Digital Hemoglobinometer	387	19,06,826
121	Table Top centrifuge with 12 tubes with stablizer	6962	41,71,33,080
122	Lab Refrigerator (400 Liters)	1973	19,63,76,261
123	Lab Refrigerator (250 Liters)	1404	11,10,66,509
124	Brain Scanner	77	10,40,40,000
125	Microscope	173	99,25,146
126	3 Part Haematology Analyzer	1352	28,39,72,500
127	Semi Automated Biochemistry Analyzer	2495	31,93,12,152
128	Thermometer	185	32,177
129	Examination Light	54	41,04,000
130	Height Measurment Scale Wall Mounted	97	2,24,460
131	Stethoscope	92	5,09,220

132	Weighing Machine Adult	92	2,50,920
133	Weighing Machine Pediatric	92	2,50,920
134	LED Torch	164	1,37,970
135	Sphygmometer	158	3,96,900
136	Otoscope	45	13,68,000
137	Tuning fork	45	54,000
138	Hammer	45	34,560
139	Tounge Depressor	45	12,960
140	Examination Table with IV Stand Foot Step	101	44,55,000
141	Revolving Stool	1710	60,87,600
142	LED X-Ray Viewing Box	97	22,44,600
143	Fetal Doppler	18	1,51,200
144	Fetal Monitor	29	65,28,600
145	Fetoscope	18	4,608
146	Infantometer	41	1,16,640
147	Stadiometer	23	80,100
148	100 M.A X-Ray Machine (Mobile)	2	17,24,400
149	ECG Machine	5	2,70,000
150	Colour Doppler Ultrasound Machine with 4 Probes: Abdomen, Pediatric	2	1,45,80,000
151	Ultra-Sonogram (obs & gyne)	2	38,70,000
152	Portable Ultrasound	2	38,70,000
153	EMG Machine	2	10,80,000
154	ECHO Machine	2	38,70,000
155	Lead Partition	11	5,85,000
156	OT Table	2	11,70,000
157	OT Light	2	9,90,000
158	Laryngoscopes with Blades	122	6,36,660
159	Crash Cart Trolley	149	83,16,000

160	Medicine Trolley	108	32,83,200
161	Surgical Scrub	2	3,87,000
162	Anesthesia Workstation	14	1,83,60,000
163	Suction Machine (Electrical)	2	57,600
164	Autorefractor Keratometry	2	28,80,000
165	Digital Lensometer	2	23,40,000
166	Non Contact Tonometer	2	39,42,000
167	Slitlamp	2	61,20,000
168	Digital Phoraptor	2	63,90,000
169	Operating Microscope	2	1,76,40,000
170	Operating Microscope Basic	2	81,00,000
171	Pheco Machine	2	1,15,20,000
172	Yag Laser	2	1,08,00,000
173	IOL Master	2	1,71,00,000
174	Crossmatch workstation	2	10,08,000
175	Centrifuge	9	17,64,000
176	Tube Sealer	2	6,66,000
177	Blood Collection Monitor	2	6,75,000
178	Doner Coach	2	8,82,000
179	Blood bank Refrigerator	2	15,57,000
180	Plasma Freezer -80 Degree	2	28,44,000
181	Plasma Extractor	2	2,70,000
182	VDRL Shakar	2	1,78,200
183	Blood Bank Centrifuge 12 Bucket	2	88,20,000
184	Composcale	2	1,80,000
185	Hand Stripper	2	1,24,200
186	Platelet Agitator with Incubator	2	15,21,000
187	Portable donar chair	2	57,600
188	Calorimeter	2	1,16,100

189	Elisa reader and washer	5	23,40,000
190	Semi Auto analyser	2	5,31,000
191	Electrolyte Analyzer	2	4,95,000
192	Reagent Refrigerator	2	4,68,000
193	Fully Automatic Biochemistry Analyser High End	2	1,60,20,000
194	Tube Lebling Analyzer	2	27,90,000
195	Advance 5 part cell counter with autoloader	2	91,80,000
196	Cell Counter	2	7,56,000
197	ESR Analyzer	2	5,94,000
198	Binocular Microscope	5	3,42,000
199	Coagulation Analyzer	2	5,31,000
200	HBA1C Analyzer fully automatic	2	57,60,000
201	Biosafety Cabinate	2	27,90,000
202	Microtom	2	53,37,000
203	Tissue processor	2	1,45,80,000
204	Embeding system	2	1,63,81,800
205	Slide stainers	2	30,33,000
206	Cytocentrifuge	2	14,40,000
207	Grossing station	2	85,77,000
208	RTPCR analyzer	2	70,20,000
209	Nucleic acid extraction system	2	91,80,000
210	Gel documentation system	2	55,80,000
211	Laminar flow hood	2	11,52,000
212	Deep freezer -20	2	20,69,280
213	Deep freezer -80	2	25,86,600
214	Chemiluminescence immunology analyzer clia	2	79,20,000
215	Incubators	5	8,62,200
216	Water Bath for Serology	2	1,89,684

217	Hot Air Oven	2	2,58,660
218	Rotor/Shaker	2	2,06,928
219	Time stop watch	2	5,166
220	Alarm Clock	2	5,166
221	HB Meter	2	58,500
222	PH Meter	2	1,29,330
223	Dental Chair with Accessoriess	18	93,60,000
224	Dental Xray Machine with RVG	7	21,06,000
225	BP Apparatus	38	96,390
226	Defibrillator	14	52,92,000
227	Video Bronchoscope	2	25,86,600
228	Audiometer	2	11,16,000
229	Tympanometer	2	11,70,000
230	Berra Machine	2	43,11,000
231	ENT Head light	2	51,732
232	OAE Screeing Machine	9	1,51,74,720
233	Moterized Labor Table	20	71,28,000
234	Infant Weighing Scale	34	3,10,500
235	Neonate Pulse Oxymeter Table Top	32	19,40,400
236	Vacuum Extractor	20	27,93,528
237	Beds with foam Mattress	1393	4,40,10,900
238	Bed side Locker Delux	1555	1,43,03,700
239	SpO2 Monitor	117	72,07,200
240	Nebulizer Ultrasonic	158	9,65,664
241	IV Stand	1469	39,96,360
242	Over Bed Table	1589	1,84,26,600
243	Motorized ICU Beds	106	2,02,61,700
244	Syringe Infusion pump	90	37,93,680
245	Multipara Monitor with Stands	65	65,00,988

246	Strecher on Trolley	54	9,93,600
247	Pediatric ICU Beds	29	17,31,600
248	Pulse Oximeter with Pediatric Probe	11	6,93,000
249	Pediatric Ventilator	7	1,55,19,600
250	ABG Machine	2	20,34,792
251	Portable LED Standing Lights	2	1,71,000
252	Foot Operated Suction Machine	45	2,75,904
253	Bilirubinometer	16	42,84,000
254	Bubble Cpap	29	1,22,85,000
255	High Flow Nasal Machine	7	29,16,000
256	Open Care Radiant Warmer	99	64,94,400
257	Phototherapy Single Surface LED	34	13,23,000
258	Transport Incubator	14	63,72,000
259	Oxygen Hood	79	1,85,850
260	Pulse Oximeter Table Top	11	6,93,000
261	Cardiac Markar	2	15,51,960
262	Glucometer	45	48,600
263	Dressing Drum small	101	3,49,191
264	Dressing Drum Medium	101	3,87,990
265	Dressing Drum Large	101	5,81,985
266	Instrument try Small	101	2,32,794
267	Instrument try Medium	101	2,71,593
268	Instrument try Large	101	3,10,392
269	NO2 Cylinder	68	7,75,980
270	CO2 Cylinder	68	7,75,980
271	Anesthesia Workstation High End with accessories	11	6,57,00,000
272	C Arm Machines flat panel	7	2,37,60,000
273	Neumatic Torniquet	2	6,12,000

274	Power Bone Drill Machine	2	35,10,000
275	Cautery Machines High End with accessories	11	1,44,90,000
276	Plasma Sterilizer with accessories	11	8,82,00,000
277	Harmonic Generator with accessories	11	5,76,00,000
278	4K Laparoscope with hand instruments and accessories	11	13,68,00,000
279	Cautery Pad with cautery machine with accessories	11	86,22,000
280	Debrither with Microdrill with accessories	11	4,27,50,000
281	Smoke Evacuator with accessories	11	2,61,00,000
282	LSCS Set of 58 Instruments	20	14,74,362
283	Laparotomy set of 83 Instrument	27	27,93,528
284	MTP Set of 15 Instrument	20	6,98,382
285	Trachestomy Set of 32 Instruments	27	51,73,200
286	Eye surgery Set of 43 Instruments	14	18,62,352
287	Abdominal Tubectomy set of 63 Instrument	20	20,95,146
288	Abdominal Hysterectomy Set	20	32,59,116
289	Vaginal Hysterectomy Set	20	32,59,116
290	STSG Set	27	22,76,208
291	Delivery kit set of 20 instruments	20	13,96,764
292	VASECTOMY SET of 2 Instrument	27	82,771
293	Post Martum Kit	14	11,38,104
294	Chalazion set of 25 Instruments	14	10,13,947
295	Ortho Instrument Set	14	18,62,352
296	ENT Set 32 instruments	9	20,69,280
297	Automated System to Treat Liquid medical Waste	11	94,50,000
298	Needle Cum Syringe Destroyer	338	12,93,300
299	Three Segment Waste Segregation Trolley	113	68,97,600

300	Sharp Container	2250	4,31,100
301	Three Bucket System for Cleaning	113	21,55,500
302	Waste Transportation Trolley	45	8,62,200
303	Autoclave 4 Drum	14	1,13,81,040
304	Three seater Chair	225	65,52,720
305	Cupboard	113	30,17,700
306	Medicine Rack	113	21,55,500
307	Instrument Cabinete	113	25,86,600
308	Bedside Screens	225	13,79,520
309	Oxygen Cylinder Trolley	23	89,669
310	Video Laryngoscope	9	14,48,496
311	Cough Assisst Machine	7	35,10,000
312	Blood Warmer	23	77,59,800

Appendix B: Checklist

(Mandatory Documents to be uploaded online in the EOI)

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 EOI Due Date.

Sr. No.	Documents	Submitted	Not Submitted
1	Legal Entity Document		
2	Manufacturer/Importer		
3	Manufacturer's Authorization (Annexure XIII)		
4	Manufacturing License		
5	DPIIT (Foreign Border)		
6	Product used in country of origin		
7	Import Export Certificate for imported equipments		
8	Affidavit of imports for last three years imported equipments		
9	Undertaking by EoI Respondent (Annexure I)		
10	Proforma for Production and sale Statement (Annexure II)		
11	Annual Turnover and positive net worth Certificate (Annexure III)		
12	Supply orders in past 3 years (Govt/State/Pvt)		
13	Non-Blacklisting affidavit (Annexure V)		
14	Production capacity		
15	Mandate Form (Annexure VI)		
16	Power of Attorney (Annexure VII)		
17	Brochure / Product Literature		
18	Annual Turnover Statement (Annexure VIII)		
19	Litigation Affidavit (Annexure IX)		
20	Delivery and Supply Capability (Annexure X)		
21	Respondent Classification Details (Annexure XI)		
22	Manufacturer's Authorisation Form (Annexure XII)		
23	Proof of details of location of Service stations		
24	GST Registration		
25	PAN/TAN		
26	MSME Certificate		
27	EM II for medium Enterprises		

28	BIS/CE/USFDA certificate		
29	CDSCO		
30	ISO 13485		
31	Installation Prerequisites		

Annexure I: Undertaking by EOI Respondent

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

Subject: EOI for the [*****]

Dear Sir,

With reference to your EOI dated _____, I, having examined the document and understood its contents, hereby submit my/our EOI for the aforesaid [*****]. The EOI is submitted voluntarily and without reservation, solely for market assessment purposes.

1. I/ We acknowledge that MMGPA will be relying on the information provided in the EOI and the documents accompanying such EOI, 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 EOI are true copies of their respective originals.
2. I/ We shall make available to MMGPA any additional information it may find necessary or require supplementing or authenticating the EOI.
3. I/ We declare that:
 - a) I/ We have examined and have no reservations to the EOI, including any Addendum / Corrigendum issued by MMGPA.
 - b) I/ We do not have any conflict of interest in accordance with the EOIs; and
 - c) I/We have not directly or indirectly or through an agent engaged or indulged in any unethical practice, as defined in the EOI, in respect of any tender or request for proposal issued by or any agreement entered into with MMGPA or any other public sector enterprise or any government, Central or State.
4. I/ We understand that this EOI is purely for market analysis purpose and you may not issue any following tender or empanelment process after this.
5. I/ We certify that we the Annual Turnover and Net Worth given by us is correct to the best of our knowledge.
6. 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.
7. 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.
8. I/ We further certify that no investigation which materially affects regulatory compliance or manufacturing/import operations is pending either against us or against our associates or against our CEO or any of our directors/ managers/ employees / Partners /Trustees.
9. 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 EOI Due Date.

10. The power of attorney for signing of EOI, as per format provided at Annexure-6 of the EOI, is also enclosed.

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

Yours faithfully,

Date:

(Signature, Name and designation of the Authorised signatory)

Place:

(Name and seal of Responder)

Annexure II: Proforma for Production And Sale Statement

(For a period of last 3 Years)

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

Sr. No.	Year	Name and full Address of the Purchaser	Purchasing Entity (Gov./Semi Gov./Private hospitals/Registered under Trust)	Name of the Product	Purchase Order No. & Date	Purchase Order Quantity	Purchase Order Value (in Rs.)	Quantity		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 with client's satisfactory certificates. All purchase orders should be enclosed in the sequence as per the data provided in table above.
2. All the data provided in the above table has been verified by undersigned CA.

Name, Membership number and signature of the Chartered Accountant:

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]

Disclaimer: If, 1. The bidder is OEM , then the details of Annexure III of production in last three years

Annexure III: Average Annual Turnover and Net Worth of the Bidder

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

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

Sr. No.	Year	Turnover (In Rs.)	Positive Net worth (Yes/ No)
1	2022-23		
2	2023-24		
3	2024-25		
4	Average Annual Turnover of above 3 years		

This is to certify that the Net worth of (*name of Bidder*) is Positive for last 3 (three) Financial Years i.e., (2022-23, 2023-24, 2024-25) as per the Audited Financial Statements.

For the purposes of this RFP, net worth (the “**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 corpus and reserve amounts as certified by the chartered accountant/statutory auditor having valid registration.

Note:

- 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.
- Provide supporting Audited Financial Statements (Balance Sheets, Profit and Loss Statements, etc.) of the bidding organization/ firm.

Name, Membership number and signature of the Chartered Accountant:

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 IV: Details of Manufacturing Unit

1. **Name of the Manufacturer:**
2. **Full address:**
3. **Phone Nos.:**
4. **Fax No.:**
5. **Email ID:**
6. **Date of inception:**
7. **License No. & date:**
8. **Issued by:**
9. **Valid up to:**
10. **RTGS (Real Time Gross Settlement) System or Core Banking A/c No.:**
11. **Details of installed production capacity for 1 year:**

Sr. No.	Equipment name	Total Production Capacity	Actual Production	Installed Quantity
1				
2				
3				
4				
5				

6				
---	--	--	--	--

Date:

Seal

Signature

Name (in capital letters)

Note: The details of manufacturing unit shall be for the premises where item quoted are actually manufactured.

THE DETAILS OF FACTORY PREMISES

Person In-charge of Factory

Name :

Phone No. :

Mobile No. :

Nearest Land mark 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 :

Name of designation of the authorized signatory

Note: The details of manufacturing unit shall be for the premises where item quoted are actually manufactured.

Annexure V: Undertaking for Non-Blacklisting

(on Letterhead of the Respondent)

EOI reference No: E-297 /MMGPA/ EOI for Equipment (2026-27)

1. I/We, (Name of the Responder), the Responder herein, 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 EOI 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

Signature of Authorized Signatory

Date

Place

Verification

I, the above named [*Name of the Responder*], 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 VI: 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

(Name of the person signing & designation)

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

Annexure VII: Format for Power of Attorney for signing of EOI

(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 EOI for the [*****] ,proposed or being developed by the [*****] (the “Purchaser”) including but not limited to signing and submission of all applications, EOIs and other documents and writings, participate in Responders' and other conferences and providing information / responses to MMGPA, representing us in all matters before MMGPA, signing and execution of all contracts including the Contract and undertakings consequent to acceptance of our EOI, and generally dealing with MMGPA in all matters in connection with or relating to or arising out of our EOI for the said Contract and/or upon award thereof to us and/or till the entering into of the Contract with MMGPA.

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____

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 Responder 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 Responder.

- 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 Responders 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 VIII: Annual Turnover Statement for Three Years

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		
2		
3		
4		
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 IX: Litigation Affidavit

(on Non-judicial Stamp Paper of Rs.500/-)

EOI reference No: E-297 /MMGPA/ EOI for Equipment (2026-27)

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 X: Delivery Acceptance Certificate

FINAL ACCEPTANCE CERTIFICATE

Date: / /2026.

To,
Chief Executive Officer,
Maharashtra Medical Goods Procurement Authority,
1st Floor, Aarogya Bhawan,
St. George's Hospital Compound, Mumbai 400001
Maha.mmga2023@gmail.com

Subject: Acceptance Certificate for the supply of

Reference: Purchase Order No.

As per above mentioned reference purchase order, the said material has been received by this office / hospital in good condition & as per terms & condition of the purchase order. The details of the same are as follows;

Sr. No.	Particular(s)	Specific information
1	Name of Material Received	
2	Quantity as per Purchase Order	
3	Invoice Number with Date	
4	Batch Number(s) with Quantity	
5	Total Quantity Received	
6	Date of Receipt of Material	
7	Entry Taken in Stock Book	
8	Stock Book No.	
9	Page Number	
10	Name of the Supplier	
11	Name of the Manufacturer	
12	Address of the Manufacturer as per Invoice	
13	Material Received as per Technical Specifications mentioned in Purchase Order (YES/NO)?	

**This is certifying that, the above information is true on the basis of office record.

Name of Pharmacist/ Store In-charge;

Name of Office/ Hospital In-charge;

(Signature with Stamp)

(Signature with Stamp)

Date:

Place:

Mail id for Acceptance: maha.mmga2023@gmail.com

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EOI for Equipments

Annexure XI: Respondent Classification Details

(To be submitted on the letterhead of the EOI Respondent)

This Annexure captures classification details for MSME status, social category, and geographical presence.

A. Basic Details

Sr. No.	Parameter	Details
1	Name of EOI Respondent	
2	Type of Entity (Company / LLP)	
3	Nature (OEM / Authorized Importer)	

B. Classification Details

Sr. No.	Parameter	Details
1	MSME Status- Trading / Manufacturer or Both Please specify (Yes/No)	
2	MSME Registration Number	
3	SC/ST Ownership (Yes/No)	
4	Certificate Details	

C. Classification

Sr. No.	Classification Parameter	Select One
1	Location Classification	☐ Within Maharashtra ☐ Outside Maharashtra
2	MSME Category	☐ MSME (Within Maharashtra) ☐ MSME (Outside Maharashtra) ☐ Non-MSME
3	SC/ST Category	☐ SC/ST (Within Maharashtra) ☐ SC/ST (Outside Maharashtra) ☐ Not Applicable

Declaration

I/We hereby declare that the information furnished above is true and correct to the best of my/our knowledge and belief.

Authorized Signatory

Name: _____

Designation: _____

Date: _____

Place: _____

Annexure-XII: Manufacturer's Authorization Form

(Manufacturer's or Producer's Letter head)

To,
The Chief Executive Officer,
Maharashtra Medical Goods Procurement Authority,
1st Floor, Arogya Bhawan,
P.D' Mello Road, Mumbai- 400001.

WHEREAS (*Name of Manufacturer or producer*) (hereinafter, "we" "us") who is established and reputable manufacturer's or producers of (*name and/or description of Goods requiring this authorization*) having production facilities at (*Insert address of the factory*) do hereby authorize (*name and address of Bidder*) (herein after, the "bidder") to submit a bid, and sign the Contract with you against Request For Proposal ref no. (*Title and reference of RFP*) including the above goods produced by us.

We hereby extend our full guarantee and warranty for the above specified Goods described above in accordance with the terms and conditions of this Request for Proposal and Contract to be executed between the Bidder and Authority.

For and on behalf of the Manufacturer or Producer

Signed: _____

Date: _____

In the capacity of (*Title, position, or other appropriate designation*) and duly authorized to issue Authorization Form on behalf of (*Name of Manufacturer or producer*)

Note:

This Letter /form should be signed by a person competent and having the power of attorney/authority to legally bind the manufacturer. This should be included by the bidder in it's bid.

This Letter /form is required to be provided by Importer.