

बिड दस्तावेज़ / Bid Document

बिड विवरण/Bid Details	
बिड बंद होने की तारीख/समय /Bid End Date/Time	01-05-2026 15:00:00
बिड खुलने की तारीख/समय /Bid Opening Date/Time	01-05-2026 15:30:00
बिड पेशकश वैधता (बंद होने की तारीख से)/Bid Offer Validity (From End Date)	180 (Days)
मंत्रालय/राज्य का नाम/Ministry/State Name	Ministry Of Health And Family Welfare
विभाग का नाम/Department Name	Department Of Health And Family Welfare
संगठन का नाम/Organisation Name	All India Institute Of Medical Sciences (aiims)
कार्यालय का नाम/Office Name	Aiims, New Delhi
कुल मात्रा/Total Quantity	1600000
वस्तु श्रेणी /Item Category	Single Use Sterile Hypodermic Syringes for Human Use Conforming to IS 10258 (Part 1) (Q2)
वर्षों के अनुभव के लिए एमएसई को छूट प्राप्त है / MSE Relaxation for Years Of Experience	Yes Complete
वर्षों के अनुभव के लिए स्टार्टअप को छूट प्राप्त है / Startup Relaxation for Years Of Experience	Yes Complete
विक्रेता से मांगे गए दस्तावेज़/Document required from seller	Experience Criteria,Certificate (Requested in ATC),OEM Authorization Certificate,Additional Doc 1 (Requested in ATC) *In case any bidder is seeking exemption from Experience / Turnover Criteria, the supporting documents to prove his eligibility for exemption must be uploaded for evaluation by the buyer
क्या आप निविदाकारों द्वारा अपलोड किए गए दस्तावेजों को निविदा में भाग लेने वाले सभी निविदाकारों को दिखाना चाहते हैं? संदर्भ मेन् है/Do you want to show documents uploaded by bidders to all bidders participated in bid?	Yes (Documents submitted as part of a clarification or representation during the tender/bid process will also be displayed to other participated bidders after log in)
बिड लगाने की समय सीमा स्वतः नहीं बढ़ाने के लिए आवश्यक बिड की संख्या। / Minimum number of bids required to disable automatic bid extension	3
दिनों की संख्या, जिनके लिए बिड लगाने की समय-सीमा बढ़ाई जाएगी। / Number of days for which Bid would be auto-extended	7

बिड विवरण/Bid Details	
ऑटो एक्सटेंशन अधिकतम कितनी बार किया जाना है। / Number of Auto Extension count	1
बिड से रिवर्स नीलामी सक्रिय किया/Bid to RA enabled	Yes
रिवर्स नीलामी योग्यता नियम/RA Qualification Rule	H1-Highest Priced Bid Elimination
बिड का प्रकार/Type of Bid	Two Packet Bid
प्राथमिक उत्पाद श्रेणी/Primary product category	Single Use Sterile Hypodermic Syringes for Human Use Conforming to IS 10258 (Part 1)
तकनीकी मूल्यांकन के दौरान तकनीकी स्पष्टीकरण हेतु अनुमत समय /Time allowed for Technical Clarifications during technical evaluation	2 Days
निरीक्षण आवश्यक (सूचीबद्ध निरीक्षण प्राधिकरण /जेम के साथ पूर्व पंजीकृत एजेंसियों द्वारा)/Inspection Required (By Empanelled Inspection Authority / Agencies pre-registered with GeM)	No
मूल्यांकन पद्धति/Evaluation Method	Item wise consignee wise evaluation
मध्यस्थता खंड/Arbitration Clause	No
सुलह खंड/Mediation Clause	No

ईएमडी विवरण/EMD Detail

एडवाइजरी बैंक/Advisory Bank	State Bank of India
Schedule 1 ईएमडी राशि/EMD Amount (In INR)	12000
Schedule 2 ईएमडी राशि/EMD Amount (In INR)	16000
Schedule 3 ईएमडी राशि/EMD Amount (In INR)	28000

ईपीबीजी विवरण /ePBG Detail

एडवाइजरी बैंक/Advisory Bank	State Bank of India
ईपीबीजी प्रतिशत (%) /ePBG Percentage(%)	3.00
ईपीबीजी की आवश्यक अवधि (माह) /Duration of ePBG required (Months).	26

(a). जेम की शर्तों के अनुसार ईएमडी छूट के इच्छुक बिडर को संबंधित कटेगरी के लिए बिड के साथ वैध समर्थित दस्तावेज प्रस्तुत करने है। एमएसई कटेगरी के अंतर्गत केवल वस्तुओं के लिए विनिर्माता तथा सेवाओं के लिए सेवा प्रदाता ईएमडी से छूट के पात्र हैं। व्यापारियों को इस नीति के दायरे से बाहर रखा गया है।/EMD EXEMPTION: The bidder seeking EMD exemption, must submit the valid supporting document for the relevant category as per GeM GTC with the bid. Under MSE category, only manufacturers for goods and Service Providers for Services are eligible for exemption from EMD. Traders are excluded from the purview of this Policy.

(b). The EMD Amount will be applicable for each schedule/group selected during Bid creation.

(c).ईएमडी और संपादन जमानत राशि, जहां यह लागू होती है, लाभार्थी के पक्ष में होनी चाहिए। / EMD & Performance security should be in favour of Beneficiary, wherever it is applicable.

लाभार्थी /Beneficiary :

Aiims Jpnatc Account
 Aiims, New Delhi, Department of Health and Family Welfare, All India Institute of Medical Sciences (AIIMS),
 Ministry of Health and Family Welfare
 (Aiims Jpnatc Account)

बोली विभाजन लागू नहीं किया गया/ Bid splitting not applied.

एमआईआई खरीद वरीयता / MII Purchase Preference

एमआईआई खरीद वरीयता / MII Purchase Preference	Yes
मेक इन इंडिया विक्रेताओं को खरीद में प्राथमिकता, यदि उनका मूल्य L1+X% तक की सीमा में है / Purchase Preference to MII sellers available upto price within L1+X%	20
मेक इन इंडिया खरीद में प्राथमिकता के लिए बिड की मात्रा का अधिकतम प्रतिशत / Maximum Percentage of Bid quantity for MII purchase preference	50
सार्वजनिक खरीद (मेक-इन-इंडिया को प्राथमिकता) आदेश 2017 के अनुसार केवल क्लास 1/क्लास 2 के स्थानीय आपूर्तिकर्ताओं को ही भागीदारी की अनुमति है दिनांक 16.09.2020 (समय-समय पर संशोधित एवं लागू) / Allow participation only from Class 1/Class 2 local suppliers as per the Public procurement(Preference to Make-in-india) order 2017 date 16.09.2020(as amended and applicable time to time)	Yes, in compliance with the MII ORDER : DPIIT Order(as amended and applicable time to time)

एमएसई खरीद वरीयता/MSE Purchase Preference

एमएसई खरीद वरीयता/MSE Purchase Preference	Yes
सूक्ष्म और लघु उद्यम मूल उपकरण निर्माताओं को खरीद में प्राथमिकता, यदि उनका मूल्य L1+X% तक की सीमा में हो / Purchase Preference to MSE OEMs available upto price within L1+X%	15
सूक्ष्म और लघु उद्यम को खरीद में प्राथमिकता के लिए बिड की मात्रा का अधिकतम प्रतिशत / Maximum Percentage of Bid quantity for MSE purchase preference	25

1. If the bidder is a Micro or Small Enterprise as per latest orders issued by Ministry of MSME, the bidder shall be relaxed from the eligibility criteria of "Experience Criteria" as defined above subject to meeting of quality and technical specifications. The bidder seeking Relaxation from Experience Criteria, shall upload the supporting documents to prove his eligibility for Relaxation.

2. If the bidder is a DPIIT registered Startup, the bidder shall be relaxed from the the eligibility criteria of "Experience Criteria" as defined above subject to their meeting of quality and technical specifications. The bidder seeking Relaxation from Experience Criteria, shall upload the supporting documents to prove his eligibility for Relaxation.

3. Preference to Make In India products (For bids < 200 Crore):Preference shall be given to Class 1 local supplier as defined in public procurement (Preference to Make in India), Order 2017 as amended from time to time and its subsequent Orders/Notifications issued by concerned Nodal Ministry for specific Goods/Products. The minimum local content to qualify as a Class 1 local supplier is denoted in the bid document. If the bidder wants to avail the Purchase preference, the bidder must upload a certificate from the OEM regarding the percentage of the local content and the details of locations at which the local value addition is made along with their bid, failing which no purchase preference shall be granted. In case the bid value is more than Rs 10 Crore, the declaration relating to percentage of local content shall be certified by the statutory auditor or cost auditor, if the OEM is a company and by a practicing cost accountant or a chartered accountant for OEMs other than companies as per the Public Procurement (preference to Make-in -India) order 2017 dated 04.06.2020. Only Class-I and Class-II Local suppliers as per MII order dated 4.6.2020 will be eligible to bid. Non - Local suppliers as per MII order dated 04.06.2020 are not eligible to participate. However, eligible micro and small enterprises will be allowed to participate .The buyers are advised to refer the OM No.F.1/4/2021-PPD dated 18.05.2023.

[OM No.1 4 2021 PPD dated 18.05.2023](#) for compliance of Concurrent application of Public Procurement Policy for Micro and Small Enterprises Order, 2012 and Public Procurement (Preference to Make in India) Order, 2017.

4. Purchase preference will be given to MSEs having valid Udyam Registration and whose credentials are validated online through Udyam Registration portal as defined in Public Procurement Policy for Micro and Small Enterprises (MSEs) Order, 2012 dated 23.03.2012 issued by Ministry of Micro, Small and Medium Enterprises and its subsequent Orders/Notifications issued by concerned Ministry. If the bidder wants to avail themselves of the Purchase preference, the bidder must be the manufacturer / OEM of the offered product on GeM. Traders are excluded from the purview of Public Procurement Policy for Micro and Small Enterprises and hence resellers offering products manufactured by some other OEM are not eligible for any purchase preference. In respect of bid for Services, the bidder must be the Service provider of the offered Service. Relevant documentary evidence in this regard shall be uploaded along with the bid in respect of the offered product or service and Buyer will decide eligibility for purchase preference based on documentary evidence submitted, while evaluating the bid. If L-1 is not an MSE and MSE Seller (s) has / have quoted price within L-1+ 15% (Selected by Buyer) of margin of purchase preference /price band defined in relevant policy, such MSE Seller shall be given opportunity to match L-1 price and contract will be awarded for 25% (selected by Buyer) percentage of total quantity. The buyers are advised to refer the OM No. F.1/4/2021-PPD dated 18.05.2023 [OM No.1 4 2021 PPD dated 18.05.2023](#) for compliance of Concurrent application of Public Procurement Policy for Micro and Small Enterprises Order, 2012 and Public Procurement (Preference to Make in India) Order, 2017. Benefits of MSE will be allowed only if seller is validated on-line in GeM profile as well as validated and approved by Buyer after evaluation of documents submitted.

5. Estimated Bid Value indicated above is being declared solely for the purpose of guidance on EMD amount and for determining the Eligibility Criteria related to Turn Over, Past Performance and Project / Past Experience etc. This has no relevance or bearing on the price to be quoted by the bidders and is also not going to have any impact on bid participation. Also this is not going to be used as a criteria in determining reasonableness of quoted prices which would be determined by the buyer based on its own assessment of reasonableness and based on competitive prices received in Bid / RA process.

6. Reverse Auction would be conducted amongst all the technically qualified bidders except the Highest quoting bidder. The technically qualified Highest Quoting bidder will not be allowed to participate in RA. However, H-1 will also be allowed to participate in RA in following cases:

- i. If number of technically qualified bidders are only 2 or 3.
- ii. If Buyer has chosen to split the bid amongst N sellers, and H1 bid is coming within N.
- iii. In case Primary product of only one OEM is left in contention for participation in RA on elimination of H-1.
- iv. If L-1 is non-MSE and H-1 is eligible MSE and H-1 price is coming within price band of 15% of Non-MSE L-1
- v. If L-1 is non-MII and H-1 is eligible MII and H-1 price is coming within price band of 20% of Non-MII L-1

मूल्यांकन विधि(मदवार मूल्यांकन विधि) / Evaluation Method (Item Wise Consignee Wise Evaluation Method)

Contract will be awarded schedulewise and the determination of L1 will be done separately for each schedule. The details of item-consignee combination covered under each schedule are as under:

मूल्यांकन अनुसूचियां / Evaluation Schedules	वस्तु/श्रेणी / Item/Category	Consignee Address	मात्रा / Quantity
Schedule 1	Single Use Sterile Hypodermic Syringes For Human Use Conforming To Is 10258 (part 1)	Safdarjang Enclave , Raj Nagar , New Delhi 110029	500000
Schedule 2	Single Use Sterile Hypodermic Syringes For Human Use Conforming To Is 10258 (part 1)	Safdarjang Enclave , Raj Nagar , New Delhi 110029	500000
Schedule 3	Single Use Sterile Hypodermic Syringes For Human Use Conforming To Is 10258 (part 1)	Safdarjang Enclave , Raj Nagar , New Delhi 110029	600000

Single Use Sterile Hypodermic Syringes For Human Use Conforming To IS 10258 (Part 1) (500000 packet)

(क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक/Minimum 50% and 25% Local Content required for qualifying as Class 1 and Class 2 Local Supplier respectively)

तकनीकी विशिष्टियाँ /Technical Specifications

* जेम केटेगरी विशिष्टि के अनुसार / As per GeM Category Specification

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
PRODUCT INFORMATION	Needle included	Yes
	Syringe type based on number of pieces	Three Piece Syringe (Barrel and Piston, plunger and plunger stopper are two separate components of different materials)
	Needle cap or shield to protect the needle	Yes
	Nominal Capacity of Syringe (ml)	< 2
PACKAGING	Number of pieces in a Pack	100
CERTIFICATIONS	Compliance to Drugs and Cosmetic Act 1940 and Medical Device Rules (MDR) 2017 as amended till date	Yes
	Availability of valid Medical Device license for the product issued from the competent authority	Yes

Additional Specification Parameters - Single Use Sterile Hypodermic Syringes For Human Use Conforming To IS 10258 (Part 1) (500000 packet)

Specification Parameter Name	Bid Requirement (Allowed Values)
Specification 1 (Specifications for Hypodermic Sterile Syringe with Needle 2mL)	Nominal capacity of syringe (ml)- 2 mL, Tolerance, dead space, scale dimensions shall be as per Table 1 of IS 10258 part 1(Latest), Syringe type based on number of pieces-Three Piece syringe, Needle if provided conforming to Indian Standard (IS 10654/ISO 7864)
Gradations should be clear as (Specifications for Sterile Hypodermic Syringe with Needle 2mL)	For 2ml: After every 0.1ml and at very 0.5ml with different size and width of marking
Specification 2 (Specifications for Sterile Hypodermic Syringe with Needle 2mL)	Needle cap or shield to protect the needle, Barrel Material-should be made up of polypropylene, Material of Plunger-should be made up of Polypropylene, Material of gasket (For 3 piece) – Synthetic rubber or thermoplastic elastomer
Sample Submission	Kindly submit 5 Nos sample to surgical Store, JPNATC (Jai Prakash Narayan Apex Trauma Centre), AIIMS, New Delhi-110029 before the last date of submission of bid.

* Bidders offering must also comply with the additional specification parameters mentioned above.

परेषिती/रिपोर्टिंग अधिकारी तथा मात्रा/Consignees/Reporting Officer and Quantity

क्र.सं./S.No.	परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer	पता/Address	डिलीवरी अनुसूची /Delivery Schedule अनुबंध प्रारम्भ होने की तारीख से दिनों की संख्या में / (In number of days from contract start days)

क्र.सं./S.N o.	परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer	पता/Address	डिलीवरी अनुसूची /Delivery Schedule अनुबंध प्रारम्भ होने की तारीख से दिनों की संख्या में /(In number of days from contract start days)		
			मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____तक पूरी कर ली जाए /Delivery to be completed by
1	Krishna Murari Singh	110029,Safdarjang Enclave , Raj Nagar , New Delhi	25000	1	15
			25000	31	60
			25000	61	90
			25000	91	120
			25000	121	150
			25000	151	180
			25000	181	210
			25000	211	240
			25000	241	270
			25000	271	300
			25000	301	330
			25000	331	360
			25000	361	390
			25000	391	420
			25000	421	450
			25000	451	480
			25000	481	510
25000	511	540			
25000	541	570			
25000	571	600			

Single Use Sterile Hypodermic Syringes For Human Use Conforming To IS 10258 (Part 1) (500000 packet)

(क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक/Minimum 50% and 25% Local Content required for qualifying as Class 1 and Class 2 Local Supplier respectively)

तकनीकी विशिष्टियाँ /Technical Specifications

[* जेम कैटेगरी विशिष्टि के अनुसार / As per GeM Category Specification](#)

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
PRODUCT INFORMATION	Needle included	Yes
	Syringe type based on number of pieces	Three Piece Syringe (Barrel and Piston, plunger and plunger stopper are two separate components of different materials)
	Needle cap or shield to protect the needle	Yes
	Nominal Capacity of Syringe (ml)	>2 - 5
PACKAGING	Number of pieces in a Pack	100
CERTIFICATIONS	Compliance to Drugs and Cosmetic Act 1940 and Medical Device Rules (MDR) 2017 as amended till date	Yes
	Availability of valid Medical Device license for the product issued from the competent authority	Yes

Additional Specification Parameters - Single Use Sterile Hypodermic Syringes For Human Use Conforming To IS 10258 (Part 1) (500000 packet)

Specification Parameter Name	Bid Requirement (Allowed Values)
specification 1 (Specifications of Sterile Hypodermic Syringe with Needle 5mL)	Nominal capacity of syringe (ml)- 5 mL, Tolerance, dead space, scale dimensions shall be as per Table 1 of IS 10258 part 1(Latest) Syringe type based on number of pieces-Three Piece syringe Needle if provided conforming to Indian Standard (IS 10654/ISO 7864)
specification 2 (Specifications of Sterile Hypodermic Syringe with Needle 5mL)	Needle cap or shield to protect the needle, Barrel Material-should be made up of polypropylene, Material of Plunger-should be made up of Polypropylene, Material of gasket (For 3 piece) – Synthetic rubber or thermoplastic elastomer
Gradations should be clear as below (Specifications of Sterile Hypodermic Syringe with Needle 5mL)	For 5ml: After every 0.1ml and every 1ml
Sample Submission	Kindly submit 5 Nos sample to surgical Store, JPNATC (Jai Prakash Narayan Apex Trauma Centre), AIIMS, New Delhi-110029 before the last date of submission of bid.

* Bidders offering must also comply with the additional specification parameters mentioned above.

परेषिती/रिपोर्टिंग अधिकारी तथा मात्रा/Consignees/Reporting Officer and Quantity

क्र.सं./S.N o.	परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer	पता/Address	डिलीवरी अनुसूची /Delivery Schedule अनुबंध प्रारम्भ होने की तारीख से दिनों की संख्या में /(In number of days from contract start days)		
1	Krishna Murari Singh	110029,Safdarjang Enclave , Raj Nagar , New Delhi	मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____तक पूरी कर ली जाए /Delivery to be completed by
			25000	1	15
			25000	31	60
			25000	61	90
			25000	91	120
			25000	121	150
			25000	151	180
			25000	181	210
			25000	211	240
			25000	241	270
			25000	271	300
			25000	301	330
			25000	331	360
			25000	361	390
			25000	391	420
			25000	421	450
			25000	451	480
			25000	481	510
			25000	511	540
			25000	541	570
25000	571	600			

Single Use Sterile Hypodermic Syringes For Human Use Conforming To IS 10258 (Part 1) (60000 packet)

(क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक/Minimum 50% and 25% Local Content required for qualifying as Class 1 and Class 2 Local Supplier respectively)

तकनीकी विशिष्टियाँ /Technical Specifications

[* जेम कटेगरी विशिष्टि के अनुसार / As per GeM Category Specification](#)

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
PRODUCT INFORMATION	Needle included	Yes
	Syringe type based on number of pieces	Three Piece Syringe (Barrel and Piston, plunger and plunger stopper are two separate components of different materials)
	Needle cap or shield to protect the needle	Yes
	Nominal Capacity of Syringe (ml)	>5 - 10
PACKAGING	Number of pieces in a Pack	50
CERTIFICATIONS	Compliance to Drugs and Cosmetic Act 1940 and Medical Device Rules (MDR) 2017 as amended till date	Yes
	Availability of valid Medical Device license for the product issued from the competent authority	Yes

Additional Specification Parameters - Single Use Sterile Hypodermic Syringes For Human Use Conforming To IS 10258 (Part 1) (600000 packet)

Specification Parameter Name	Bid Requirement (Allowed Values)
specification 1 (Specifications for Sterile Hypodermic Syringe with Needle 10mL)	Nominal capacity of syringe (ml)- 10 mL, Tolerance, dead space, scale dimensions shall be as per Table 1 of IS 10258 part 1(Latest) Syringe type based on number of pieces-Three Piece syringe Needle if provided conforming to Indian Standard (IS 10654/ISO 7864)
specification 2 (Specifications for Sterile Hypodermic Syringe with Needle 10mL)	Needle cap or shield to protect the needle, Barrel Material-should be made up of polypropylene, Material of Plunger-should be made up of Polypropylene, Material of gasket (For 3 piece) – Synthetic rubber or thermoplastic elastomer
Gradations should be clear as below (Specifications for Sterile Hypodermic Syringe with Needle 10mL)	For 10ml syringe: After every 0.2 ml and after every 1ml
Sample Submission	Kindly submit 5 Nos sample to surgical Store, JPNATC (Jai Prakash Narayan Apex Trauma Centre), AIIMS, New Delhi-110029 before the last date of submission of bid.

* Bidders offering must also comply with the additional specification parameters mentioned above.

परेषिती/रिपोर्टिंग अधिकारी तथा मात्रा/Consignees/Reporting Officer and Quantity

क्र.सं./S.N o.	परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer	पता/Address	डिलीवरी अनुसूची /Delivery Schedule अनुबंध प्रारंभ होने की तारीख से दिनों की संख्या में /(In number of days from contract start days)		
			मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____तक पूरी कर ली जाए /Delivery to be completed by
1	Krishna Murari Singh	110029,Safdarjang Enclave , Raj Nagar , New Delhi	30000	1	15
			30000	31	60
			30000	61	90
			30000	91	120
			30000	121	150
			30000	151	180
			30000	181	210
			30000	211	240
			30000	241	270
			30000	271	300
			30000	301	330
			30000	331	360
			30000	361	390
			30000	391	420
			30000	421	450
			30000	451	480
			30000	481	510
30000	511	540			
30000	541	570			
30000	571	600			

Special terms and conditions-Version:1 effective from 29-09-2025 for category Single Use Sterile Hypodermic Syringes for Human Use Conforming to IS 10258 (Part 1)

1. All Provisions of Drugs and Cosmetics Act, 1940 and Rules (including Medical Device Rule 2017) made there under and all subsequent amendments till date will always be applicable. This will include all notifications issued by Central Drugs Standard Control Organization (CDSCO), Ministry of Health & Family Welfare (MoHFW) and Department of Pharmaceuticals (DOP), Ministry of Chemicals & Fertilizers in this regard.
2. The seller's registration and product upload on GeM is based on their declaration of possessing a valid Medical Device License for the product, certifications, test reports as per MDR (2017), latest

amended. However, buyers must check, validate and verify the details at their end for all applicable licenses and certifications e.g., validity and authenticity/genuineness of medical device license (or registration), product certifications, manufacturer's certifications/ licenses, test reports at the time of supply.

3. In case of authorized resellers/distributors, it will be the legal and regulatory liability of the manufacturer to ensure that their resellers/distributors are operating in compliance with all relevant laws and regulations, and are licensed to sell the manufacturer's products, including verifying the validity and authenticity of license held by them.
4. Any other Terms and Conditions which is not included or at variance with the conditions specified in STC/GTC, may be added by the buyer through Additional Terms and Conditions (ATC) in the bid to ensure items are procured from authentic/validated source with appropriate and applicable quality. The above terms and conditions are in reverse order of precedence i.e. BID ATC shall supersede specific STC which shall supersede General Terms and Conditions (GTC), whenever there are any conflicting provisions.
5. **Packing and Marking:** Should be as per MDR.
6. Presently, the products under this category are not covered under mandatory BIS licensing scheme. Hence, both ISI Marked and non-ISI Marked products may be available in this category. However, if a buyer intends to procure ISI Marked products, buyer may, at his discretion, opt for the same by indicating the requirement in the bid, after taking into account the number of valid licenses issued under the standard. In such cases, buyer shall verify valid BIS License as per the applicable governing standard at the time of evaluation of bid and check the ISI marking on the products at the time of receipt of material before issuing of CRAC.
The valid BIS license status of sellers may be viewed by buyer from the below link.

<https://www.manakonline.in/MANAK/ApplicationLicenceRelatedrpt>

or

https://www.services.bis.gov.in/php/BIS_2.0/bisconnect/knowyourstandards/indian_standards/isdetails

क्रेता द्वारा जोड़ी गई बिड की विशेष शर्तें/**Buyer Added Bid Specific Terms and Conditions**

1. Experience Certificate for the supply of the same to any Govt/ PSU/ any renowned private organisation along with Supply/ Purchase Order.
2. If the agency is registered under MSME or NSIC, then EMD exemption certificate needs to be enclosed.
3. Make in india specific authorisation certificate needs to be enclosed.
4. **Generic**

OPTION CLAUSE: The Purchaser reserves the right to increase or decrease the quantity to be ordered up to 50 percent of bid quantity at the time of placement of contract. The purchaser also reserves the right to increase the ordered quantity up to 50% of the contracted quantity during the currency of the contract at the contracted rates. The delivery period of quantity shall commence from the last date of original delivery order and in cases where option clause is exercised during the extended delivery period the additional time shall commence from the last date of extended delivery period. The additional delivery time shall be $(\text{Increased quantity} \div \text{Original quantity}) \times \text{Original delivery period (in days)}$, subject to minimum of 30 days. If the original delivery period is less than 30 days, the additional time equals the original delivery period. The Purchaser may extend this calculated delivery duration up to the original delivery period while exercising the option clause. Bidders must comply with these terms.

5. **Buyer Added Bid Specific ATC**

Buyer Added text based ATC clauses

Risk Purchase Clause

AIIMS is committed to ensuring uninterrupted patient care and maintaining the quality and timely availability of all the patient necessary medical items. To safeguard this objective, AIIMS reserves the right to invoke the Risk Purchase Clause in case of non-performance or default by the selected bidder (L-1). Subject to the provision under Force Majeure clause, failure to adhere to the prescribed schedule may result in penalties or invocation of the Risk Purchase Clause whereby the purchaser reserves the right to procure undelivered items from an alternate source at the defaulting bidder's risk and cost.

Purchaser's Right to accept any bid and to reject any or all bids- The purchaser reserves the right to accept

t in part or in full any bid or reject any or more bid(s) without assigning any reason or to cancel the bid process and reject all bids at any time prior to award of contract, without incurring any liability, whatsoever to the affected bidder(s).

Note* Vendors found supplying substandard products will be blacklisted and stringent penalty clause imposed.

Bidders must upload the following documents in the Technical Bid:

1. Manufacturers Authorization Certificate
2. Certification of Local Content (MII)
3. Product Compliance sheet mentioning compliance to specifications and additional specifications mentioned in bid document
4. Product Certifications
5. Non-blacklisting undertaking of OEM and distributor both
6. Bidders Registration Certificate, GST certificate, CDSCO Manufacturing drug/Medical Device license of OEM
7. Compliance to all terms and conditions of Bid
8. EMD- Rs. 12000-syringe 2ml, Rs. 16000-syringe 5ml and Rs. 28000-syringe 10ml (should also submit hard copy of EMD Receipt to Store Section (JPNATC) Basement-1, J PNATC, Near Raj Nagar & Safdarjung Hospital, AIIMS, New Delhi-110029)
9. Every item should be stamped AIIMS SUPPLY NOT FOR SALE

अस्वीकरण/**Disclaimer**

The Additional Terms and Conditions (ATC) have been incorporated by the Buyer after approval of their Competent Authority. The Buyer, is solely responsible for the impact of these clauses on the bidding process, its outcome, and consequences thereof including any restriction arising in the bidding process due to these ATCs and including the modification of technical specifications and / or terms and conditions governing the bid. All representations / grievances pertaining to the ATC clauses shall be raised with the buyer organization directly and not with GeM. If any of the clause(s) is/are incorporated by the Buyer regarding the following, the bid & resultant contract shall be treated as null & void. Further, GeM reserves the right, at its sole discretion, to cancel the bid forthwith, without issuance of any prior notice or intimation :-

1. Publishing Custom / BOQ bids for items for which regular GeM categories are available (unless such Custom / BOQ item is bunched with the major regular product Category Item).
2. Mandating procurement of / from specific Brand / Make / Model / Manufacturer / Dealer except in case of Single Bid / Proprietary Article Certificate (PAC) Buying.
3. Inclusion of disqualification criteria related to suspension of seller / service provider, where such suspension period has already expired.
4. Mandating submission of documents in physical form as a pre-requisite to qualify bidders.
5. Publishing bids on GeM for procurement of works.
6. Procurement of Goods by creating a Service bid on GeM & vice-versa.
7. Seeking sample with bid or approval of samples during bid evaluation process. However, trial / sample, as the case may be, shall be permitted in cases where trial / sample are allowed as per approved and published procurement policy of the Buyers' controlling Ministry / Department / State / Public Sector Enterprises Headquarters. If there is any violation of trial / sample clause with regard to approved policy of the Buyers' Ministry / Department / State / Public Sector Enterprises Headquarters, then this is to be determined and redressed by the concerned Buyer Organisation only.
8. Seeking experience from specific organization / department / institute only or from foreign / export experience.
9. Creating bid for items from incorrect categories.

10. Reference of conditions published on any external site or reference to external documents/clauses.
11. Asking for any Tender fee / Bid Participation fee, as the case may be.
12. Buyer added ATC Clauses which are in contravention of clauses defined in bid detail section, including specifications, EMD Detail, ePBG Detail and MII and MSE Purchase Preference sections of the bid, unless otherwise allowed by the applicable GeM GTC.
13. Any ATC clause in contravention with GeM GTC Clause 4 (xiii) (h) will be invalid. In case of multiple L1 bidders against a service bid, the buyer shall place the Contract by selection of a bidder amongst the L-1 bidders through a Random Algorithm executed by GeM system.
14. In a category based bid, adding additional items, through buyer added, additional scope of work/ additional terms and conditions/or any other document. If buyer needs more items along with the main item, the same must be added through bunching category based items or by bunching custom catalogues or bunching a BoQ with the main category based item, the same must not be done through ATC or Scope of Work.

Further, if any seller has any objection/grievance against these additional clauses or otherwise on any aspect of this bid, they can raise their representation against the same by using the Representation window provided in the bid details field in Seller dashboard after logging in as a seller. Buyer is duty bound to reply to all such representations and would not be allowed to open bids if he fails to reply to such representations.

All GeM Sellers/Service Providers shall ensure full compliance with all applicable labour laws, including the provisions, rules, schemes and guidelines under the four Labour Codes i.e. the Code on Wages, 2019; the Industrial Relations Code, 2020; the Occupational Safety, Health and Working Conditions Code, 2020; and the Code on Social Security, 2020 as and when notified and brought into force by the Government of India.

For all provisions of the Labour Codes that are pending operationalisation through rules, schemes or notifications, the corresponding provisions of the pre-existing labour enactments (such as The Minimum Wages Act, 1948, The Payment of Wages Act, 1936, The Payment of Bonus Act, 1965, The Equal Remuneration Act, 1976, The Payment of Gratuity Act, 1972, etc. and relevant State Rules) shall continue to remain applicable.

The Seller/ Service Providers shall, therefore, be responsible for ensuring compliance under:

- **All notified and enforceable provisions of the new Labour Codes as mentioned hereinabove; and**
- **All operative provisions of the erstwhile Labour Laws until their complete substitution.**

All obligations relating to wages, social security, safety, working conditions, industrial relations etc. and any other statutory requirements shall be strictly met by the Seller/ Service Provider. Any non-compliance shall constitute a breach of the contract and shall entitle the Buyer to take appropriate action in accordance with the contract and applicable law.

This Bid is governed by the General Terms and Conditions, conditions stipulated in Bid and Service Level Agreement specific to the Service, as the case may be, as provided in the Marketplace.

However, in case of Service, if any condition specified in General Terms and Conditions is contradicted by the conditions stipulated in Service Level Agreement specific to said Service, then it will over-ride the conditions in the General Terms and Conditions.

[यह बिड सामान्य शर्तों के अंतर्गत भी शासित है /This Bid is also governed by the General Terms and Conditions](#)

जेम की सामान्य शर्तों के खंड 26 के संदर्भ में भारत के साथ भूमि सीमा साझा करने वाले देश के बिडर से खरीद पर प्रतिबंध के संबंध में भारत के साथ भूमि सीमा साझा करने वाले देश का कोई भी बिडर इस निविदा में बिड देने के लिए तभी पात्र होगा जब वह बिड देने वाला सक्षम प्राधिकारी के पास पंजीकृत हो। बिड में भाग लेते समय बिडर को इसका अनुपालन करना होगा और कोई भी गलत घोषणा किए जाने व इसका अनुपालन न करने पर अनुबंध को तत्काल समाप्त करने और कानून के अनुसार आगे की कानूनी कार्यवाई का आधार होगा।/In terms of GeM GTC clause 26 regarding Restrictions on procurement from a bidder of a country which shares a land border with India, 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. While participating in bid, Bidder has to undertake compliance of this and any false declaration and non-compliance of this would be a ground for immediate termination of the contract and further legal action in accordance with the laws.

---धन्यवाद/Thank You---

