



Bid Number/बोली क्रमांक (बिड संख्या):
GEM/2024/B/5363152
Dated/दिनांक : 05-09-2024

Bid Document/ बिड दस्तावेज़

Bid Details/बिड विवरण	
Bid End Date/Time/बिड बंद होने की तारीख/समय	26-09-2024 14:00:00
Bid Opening Date/Time/बिड खुलने की तारीख/समय	26-09-2024 14:30:00
Bid Offer Validity (From End Date)/बिड पेशकश वैधता (बंद होने की तारीख से)	180 (Days)
Ministry/State Name/मंत्रालय/राज्य का नाम	Ministry Of Health And Family Welfare
Department Name/विभाग का नाम	Department Of Health And Family Welfare
Organisation Name/संगठन का नाम	All India Institute Of Medical Sciences (aiims)
Office Name/कार्यालय का नाम	Aiims, New Delhi
Total Quantity/कुल मात्रा	250000
Item Category/मद केटेगरी	Intravenous Cannulas as per IS 10555 - 5 (Q2)
MSE Exemption for Years of Experience and Turnover/ अनुभव के वर्षों से एमएसई छूट	No
Startup Exemption for Years of Experience and Turnover/ अनुभव के वर्षों से स्टार्टअप छूट	No
Document required from seller/विक्रेता से मांगे गए दस्तावेज़	Certificate (Requested in ATC), OEM Authorization Certificate *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
Do you want to show documents uploaded by bidders to all bidders participated in bid?/	Yes
Bid to RA enabled/बिड से रिवर्स नीलामी सक्रिय किया	Yes
RA Qualification Rule	50% Lowest Priced Technically Qualified Bidders
Type of Bid/बिड का प्रकार	Two Packet Bid
Time allowed for Technical Clarifications during technical evaluation/तकनीकी मूल्यांकन के दौरान तकनीकी स्पष्टीकरण हेतु अनुमत समय	2 Days
Inspection Required (By Empanelled Inspection Authority / Agencies pre-registered with GeM)	No

Bid Details/बिड विवरण**Evaluation Method/मूल्यांकन पद्धति**

Total value wise evaluation

EMD Detail/ईएमडी विवरण

Required/आवश्यकता

No

ePBG Detail/ईपीबीजी विवरण

Advisory Bank/एडवाइजरी बैंक

State Bank of India

ePBG Percentage(%) / ईपीबीजी प्रतिशत (%)

3.00

Duration of ePBG required (Months) / ईपीबीजी की अपेक्षित अवधि (महीने).

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

Beneficiary/लाभार्थी :

Director

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

MII Purchase Preference/एमआईआई खरीद वरीयता

MII Purchase Preference/एमआईआई खरीद वरीयता

Yes

MSE Purchase Preference/एमएसई खरीद वरीयता

MSE Purchase Preference/एमएसई खरीद वरीयता

Yes

1. 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.

2. 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.

3. Reverse Auction would be conducted amongst first 50% of the technically qualified bidders arranged in the order of prices from lowest to highest. Number of sellers eligible for participating in RA would be rounded off to next higher integer value if number of technically qualified bidders is odd (e.g. if 7 bids are technically qualified, then RA will be conducted amongst L-1 to L-4). In case number of technically qualified bidders are 2 or 3, RA will be between all without any elimination. If Buyer has chosen to split the bid amongst N sellers, then minimum N sellers would be taken to RA round. In case Primary products of only one OEM are left in contention for participation in RA based on lowest 50% bidders qualifying for RA, the number of sellers qualifying for RA would be increased to get at least products of one more OEM (directly participated or through its reseller) if available. Further, if bid(s) of any seller(s) eligible for MSE preference is / are coming within price band of 15% of Non MSE L-1 or if bid of any seller(s) eligible for Make in India preference is / are coming within price band of 20% of non MII L-1, then such MSE / Make in India seller shall also be allowed to participate in the RA process.

Intravenous Cannulas As Per IS 10555 - 5 (250000 pieces)

(Minimum 50% and 20% Local Content required for qualifying as Class 1 and Class 2 Local Supplier respectively/क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक)

Technical Specifications/तकनीकी विशिष्टियाँ

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

Specification	Specification Name/विशिष्टि का नाम	Bid Requirement/बिड के लिए आवश्यक (Allowed Values)/अनुमत मूल्य
Performance Parameters	Conforming standards for the I V Canulla	IS / ISO 10555-5
	Whether IV Canulla is with safety features for preventing needle stick injuries	Yes, No
	Needle Point Finish	Long bevel cut
	Needle Hub	Complying with ISO 594-1
	Needle hub fitting with 6% luer Taper	Yes
	Injection Port	With

Specification	Specification Name/विशिष्टि का नाम	Bid Requirement/बिड के लिए आवश्यक (Allowed Values)/अनुमत मूल्य
	Wings	With
	Luer Lock Plug/Cap	Yes
	Radio opaque line feature	Yes, No
Dimensional and Material Parameters	Material of Catheter	Polytetrafluoroethylene (PTFE)
	Nominal outside Diameter of catheter tube and color coding as per IS/ISO 10555-5	Orange - 14G, Medium Grey - 16G, Deep Green - 18G, Pink - 20G, Deep Blue - 22G

Additional Specification Parameters - Intravenous Cannulas As Per IS 10555 - 5 (250000 pieces)

Specification Parameter Name	Bid Requirement (Allowed Values)
SAMPLE CLAUSE	Kindly submit 10 nos. of samples on Monday to Friday at 2.00 PM to 3.00 PM to Mr. Satish Chandra, Surgical Store-1 of Main Hospital Store, AIIMS, New Delhi - 110029 before bid opening date. In case sample is not submitted, the offer shall be out rightly rejected.
SPECIFICATION	Quoted item must be comply with the specifications attached in Buyer Added Bid Specific ATC.
RISK PURCHASE	AIIMS reserves the right to impose risk purchase clause to meet the urgent patient care as "For delay in execution/supply we shall have every right to procure the same from Open Market and the additional cost of the same (Incidental/Consequential) will be recovered from the payment of the vendor dues under this risk purchase clause.
QUANTITY AND SIZE OF CANNULA	1. 50000 of 14 G 2. 50000 of 16 G 3. 50000 of 18 G 4. 50000 of 20 G 5. 50000 of 22 G
SUPPLY CLAUSE	Please provide the copy of the Supply order with Challan in triplicate at the time of Supply. Each item should be stamped with permanent Ink/Print as "AIIMS SUPPLY, NOT FOR SALE". Submit test/batch report (if applicable) along with every supply.

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

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

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

S.No./क्र. सं.	Consignee Reporting/Officer/ परेषिती/रिपोर्टिंग अधिकारी	Address/पता	Delivery Schedule/डिलीवरी अनुसूची (In number of days from contract start days/अनुबंध प्रारंभ होने की तारीख से दिनों की संख्या में)		
			Quantity/मात्रा	Delivery to start after/प्रारंभ होने की तारीख से डिलीवरी	Delivery to be completed by/डिलीवरी तक पूरी कर ली जाए
1	Satish Chandra	110029,Ansari Nagar	30000	1	15
			20000	16	30
			20000	31	60
			20000	61	90
			20000	91	120
			20000	121	150
			20000	151	180
			20000	181	210
			20000	211	240
			20000	241	270
			20000	271	300
			20000	301	360

Special terms and conditions-Version:3 effective from 25-11-2022-Version:1 effective from 04-05-2020 for category Intravenous Cannulas as per IS 10555 - 5

- All Provisions of Drugs and Cosmetics Act, 1940 and Rules made there under as amended till date will always be applicable. This will include all notifications issued by *Central Drugs Standard Control Organisation (CDSCO)*, Ministry of Health & Family Welfare (MoHFW) and Department of Pharmaceuticals (DOP), Ministry of Chemicals & Fertilizers time to time in this regard.
 - The sellers are registered on GeM based on self-declaration of valid Drug License, product certification, test reports etc. However, buyers must check and validate the details at their end for all applicable licenses and certifications e.g., validity and authenticity/genuineness of drug license, product certification, manufacturer certification/licenses, test reports etc.
 - The price offered by the seller shall not, in any case exceed the DPCO controlled price, if any, fixed by the Central/State Government, the Maximum Retail Price (MRP) and the selling price. The seller must reduce the prices if there is any reduction in DPCO ceiling price, if any.
 - 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 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., ATC shall supersede specific STC which shall supersede General Terms and Conditions ("GTC"), whenever there are any conflicting provisions.
- Special Terms and Conditions for Medical Devices and Consumables covered under Provisions of Drug and Cosmetic Act
 - For items wherever Drug Licence requirements are applicable all provisions of Drug and Cosmetic Act 1940 as amended up to date and Rules made there under will be applicable in addition to any other terms and conditions specified in the Portal.

2. Drug License: For indigenous products offered in the market, Manufacturer should have valid Drug License as per Drugs and Cosmetic Act 1940 issued by concerned State Drug Control authorities .The Seller if different from the manufacturer shall also be required to be holding Drug License for sale . In case of imported products Manufacturer shall be registered under Form no 10 with Central Drug Authorities (CDSCO) and the Seller offering imported products should be also holding valid Sales License issued by the local drug authorities. For imported products, certificate from the OEM that product is being used in the Country of Origin should be available with the Seller. It shall be the responsibility of the Seller to ensure that that the Drug License is valid for the product offered and due to any reason the drug control authorities have cancelled or suspended Drug License or convicted the manufacturer or Seller for any offence under the provisions of Drug and Cosmetic Act, Seller should immediately withdraw the product and also intimate the Buyers in case of pending orders for supplies as well as the GeM administration regarding the matter.
3. Manufacturing & Marketing Experience: Sellers offering the Products in the Portal either as Manufacturers or as Authorised Seller shall ensure that the Products offered are being Manufactured and Marketed in the country (for Indigenous Products) and Marketed (for Imported Products) continuously at least for the last 2 years
4. Certifications: Manufacturers of offered product (Offered by Manufacturers or by Authorized Seller) should be holding valid Good Manufacturing Practices Certificate (GMP) as per revised Schedule-M of Drug and Cosmetic Act 1940 as amended up to date or WHO-GMP as per norms amended up to date issued by the Licensing Authority or certificate which is at par with WHO-GMP issued by the authorities of exporting countries / COPP certificate .
5. Non Conviction Certificate: Sellers either Manufacturers or Authorized Sellers are required to ensure that they are not under conviction in terms of the provisions of Drugs & Cosmetic Act and any other law applicable in relation to the same . In case at any point of time, the Manufacturer or Authorized Seller is convicted under provisions of Drug and Cosmetic Act, it shall be their responsibility to withdraw the product immediately from the market.
6. Banning and Blacklisting: Seller either Manufacturer or Authorized Seller shall ensure that there is no banning or black listing applicable against them for the product offered on the portal due to quality failure and /or fraudulent/illegal practices or for any other reasons
7. It shall be the responsibility of the Seller either Manufacturer or Authorized Seller to ensure that manufacturer is having own in-house testing lab to carry out all the required tests as per specification and provisions of drug act as amended up to date for the quoted product and shall also forward the copies of the in-house test reports for each batch along with the supplies. For imported products , certificate from OEM regarding availability of all test facilities in house with them should be available with Seller .
8. Each lot of supplies shall be dispatched under Self Certification scheme duly supported by in house test reports. Consignees shall be at liberty to draw control Samples and send it to approved Laboratories for testing and in case of any failure , entire responsibility shall rest with Seller in addition to any penalties under the provisions of Drug Act including removal of Goods from the Consignee place. .Further administrative actions as per terms and conditions Gem Portal shall also be applicable.
9. Packing shall be as per relevant clause of Standard Specifications applicable as indicated in the Catalogue Parameters indicated in the Portal and as per provisions of Drug & Cosmetic Act as amended up to date.
10. Expiry Date: All supplies must indicate the Month of Manufacture and Expiry. In addition all supplies shall have a remaining shelf life of at least 5/6th of the stipulated shelf life at the time of delivery.
11. Recalls: If any batch is to be recalled because of problems with product quality or adverse reaction Seller will be responsible to notify the Buyer full details about the reason leading to the recall and shall take steps to replace the product in question at its own cost with a fresh batch of acceptable quality or give a refund of the value of the goods

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

1. If the agency is registered under MSME or NSIC, then EMD exemption certificate needs to be enclosed.
2. Make in india specific authorisation certificate needs to be enclosed.
3. **Buyer Added Bid Specific ATC**

Buyer uploaded ATC document [Click here to view the file.](#)

Disclaimer/अस्वीकरण

The additional terms and conditions have been incorporated by the Buyer after approval of the Competent Authority in Buyer Organization, whereby Buyer organization is solely responsible for the impact of these clauses on the bidding process, its outcome, and consequences thereof including any eccentricity / restriction arising in the bidding process due to these ATCs and due to modification of technical specifications and / or terms and conditions governing the bid. If any clause(s) is / are incorporated by the Buyer regarding following, the bid and resultant contracts shall be treated as null and void and such bids may be cancelled by GeM at any stage of bidding process without any notice:-

1. Definition of Class I and Class II suppliers in the bid not in line with the extant Order / Office Memorandum issued by DPIIT in this regard.
2. Seeking EMD submission from bidder(s), including via Additional Terms & Conditions, in contravention to exemption provided to such sellers under GeM GTC.
3. Publishing Custom / BOQ bids for items for which regular GeM categories are available without any Category item bunched with it.
4. Creating BoQ bid for single item.
5. Mentioning specific Brand or Make or Model or Manufacturer or Dealer name.
6. Mandating submission of documents in physical form as a pre-requisite to qualify bidders.
7. Floating / creation of work contracts as Custom Bids in Services.
8. Seeking sample with bid or approval of samples during bid evaluation process. (However, in bids for [attached categories](#), trials are allowed as per approved procurement policy of the buyer nodal Ministries)
9. Mandating foreign / international certifications even in case of existence of Indian Standards without specifying equivalent Indian Certification / standards.
10. Seeking experience from specific organization / department / institute only or from foreign / export experience.
11. Creating bid for items from irrelevant categories.
12. Incorporating any clause against the MSME policy and Preference to Make in India Policy.
13. Reference of conditions published on any external site or reference to external documents/clauses.
14. Asking for any Tender fee / Bid Participation fee / Auction fee in case of Bids / Forward Auction, as the case may be.

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 within 4 days of bid publication on GeM. 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.

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

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./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।

---Thank You/धन्यवाद---