बिड संख्या/Bid Number: GEM/2026/B/7109207

दिनांक /Dated: 13-01-2026

बिड दस्तावेज़ / Bid Document

बिड विवरण/Bid Details	
बिड बंद होने की तारीख/समय /Bid End Date/Time	04-02-2026 19:00:00
बिड खुलने की तारीख/समय /Bid Opening Date/Time	04-02-2026 19:30:00
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	1800000
वस्तु श्रेणी /Item Category	Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile and Single Use) Conforming to IS/ISO 10555 (Part 5) (Q2)
वर्षों के अनुभव एवं टर्नओवर से एमएसई को छूट प्राप्त है / MSE Relaxation for Years Of Experience and Turnover	Yes | Complete
स्टार्टअप के लिए अनुभव के वर्षों और टर्नओवर से छूट प्रदान की गई है / Startup Relaxation for Years Of Experience and Turnover	Yes | Complete
विक्रेता से मांगे गए दस्तावेज़/Document required from seller	Experience Criteria,Bidder Turnover,Certificate (Requested in ATC),OEM Authorization Certificate,Compliance of BoQ specification and supporting document *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 दस्तावेज़ों को निविदा में भाग लेने वाले सभी निविदाकारों को दिखाना चाहते हैं? संदर्भ मेनू 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	1

बिड विवरण/Bid Details	
दिनों की संख्या, जिनके लिए बिड लगाने की समय-सीमा बढ़ाई जाएगी। / Number of days for which Bid would be auto-extended	7
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
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
अनुमानित बिड मूल्य /Estimated Bid Value	9000000
मूल्यांकन पद्धति/Evaluation Method	Total value wise evaluation
मध्यस्थता खंड/Arbitration Clause	No
सुलह खंड/Mediation Clause	No

ईएमडी विवरण/EMD Detail

एडवाईजरी बैंक/Advisory Bank	State Bank of India
ईएमडी राशि/EMD Amount	450000

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 केटेगरी के लिए बिड के साथ वैध समर्थित दस्तावेज़ प्रस्तुत करने है। एमएसई केटेगरी 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).ईएमडी और संपादन जमानत राशि, जहां यह लागू होती है, लाभार्थी के पक्ष में होनी चाहिए। / EMD & Performance securityshould be

in favour of Beneficiary, wherever it is applicable.

लाभार्थी /Beneficiary :
AIIMS
Aiims, New Delhi, Department of Health and Family Welfare, All India Institute of Medical Sciences (AIIMS), Ministry of Health and Family Welfare
(Aiims Main Grant)

UIN Number NCTGC2415P

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 availabele 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 Micro or Small Enterprise (MSE) as per latest orders issued by Ministry of MSME, the bidder

shall be relaxed from the eligibility criteria of "Bidder Turnover" as defined above subject to meeting of quality and technical specifications. If the bidder itself is MSE OEM of the offered products, it would be relaxed from the "OEM Average Turnover" criteria also subject to meeting of quality and technical specifications. The bidder seeking Relaxation from Turnover, shall upload the supporting documents to prove his eligibility for Relaxation.
3. 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.
4. If the bidder is a DPIIT registered Startup, the bidder shall be relaxed from the the eligibility criteria of "Bidder Turnover" as defined above subject to their meeting of quality and technical specifications. If the bidder is DPIIT Registered OEM of the offered products, it would be relaxed from the "OEM Average Turnover" criteria also subject to meeting of quality and technical specifications. The bidder seeking Relaxation from Turnover shall upload the supporting documents to prove his eligibility for Relaxation.
5. 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.

6. 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.
7. 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.

8. 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

Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile And Single Use)

Conforming To IS/ISO 10555 (Part 5) (1800000 packet)

(क्रमशः श्रेणी 1 और श्रेणी 2 के स्थानीय आपूर्तिकर्ता के रूप में अर्हता प्राप्त करने के लिए आवश्यक/Minimum 50% and 20% 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	**Number of Catheter Lumen**	Single Lumen, Two Lumen, Triple Lumen
	Nominal outside diameter (mm)	0.6 mm, 0.7 mm, 0.8 mm, 0.9 mm, 1.0 mm, 1.1 mm, 1.2 mm, 1.3 mm, 1.4 mm, 1.5 mm, 1.6 mm, 1.7 mm, 1.8 mm, 1.9 mm, 2.0 mm, 2.1 mm, 2.2 mm, 2.3 mm, 2.4 mm, 2.5 mm, 2.6 mm, 2.7 mm, 2.8 mm, 3.3 mm, 3.4 mm
	Color coding (as per Table 1 of IS/ISO 10555-5)	Orange, Medium Grey, Deep Green, Pink, Deep Blue, Yellow, Violet, White, Red, Pale Blue, Light Brown
PACKAGING	**Number of pieces in a Pack**	50, 100
SHELF LIFE	Residual shelf life of the product	Agree to provide residual shelf life of atleast 75% of total shelf life at the time of delivery to the consignee
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 - Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile And Single Use) Conforming To IS/ISO 10555 (Part 5) (1800000 packet)

Specification Parameter Name	Bid Requirement (Allowed Values)
Additional Specification Parameters	All bidders must quote the product strictly as per the SPECIFICATIONS mentioned at ‘Buyer Added Bid Specific ATC’ under 'BUYER ADDED BID SPECIFIC TERMS AND CONDITIONS' in bid document. Products not conforming to the detailed specifications prescribed by the Buyer’s organization shall be summarily rejected without any further communication or opportunity for correction. The Buyer’s decision regarding conformity to the prescribed specifications shall be final and binding on all bidders.
Buyer-added Bid Specific Additional Terms and Conditions	Bidders must carefully review the enclosed document under “Buyer Added Bid Specific ATC” to fully understand all bid terms, conditions. Tenders not conforming to the prescribed T&C by the Buyer’s organization shall be summarily rejected without any further communication or opportunity for correction. The Buyer’s decision regarding conformity to the prescribed bid conditions shall be final and binding on all bidders.

Specification Parameter Name	Bid Requirement (Allowed Values)
Undertaking by Bidders	All bidders must submit a duly signed undertaking confirming full compliance with all the technical specifications and all T & C specified under the Buyer Added Bid Specific ATC of the bid document. Submission of this declaration is a mandatory requirement and Bids submitted without the required undertaking shall be summarily rejected and treated as non responsive. No correspondence, clarification, or request for reconsideration shall be entertained in this regard.
Sample Clause and Delivery Address	Bidders should submit atleast 05 number of (FIVE) samples IN EACH SIZE of the quoted product ON or BEFORE BID END DATE to Surgical Store-1, Hospital Stores, AIIMS, New Delhi-110 029. These samples will be evaluated based on actual usage and the technical requirements set by the buyer, as the product is intended for patient care services. Failure to submit samples ON or BEFORE BID END DATE will result in disqualification of the bid.
Risk Purchase Clause	AIIMS is committed to ensuring uninterrupted patient care by maintaining the quality and timely availability of all essential medical items. To uphold this objective, AIIMS reserves the right to invoke the Risk Purchase Clause in the event of non-performance or default by the selected bidder (L-1). Bidders are strongly advised to carefully review ATC, Point No. 08 for for detailed provisions related to the Risk Purchase Clause before quoting their 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	मात्रा /Quantity	डिलीवरी के दिन/Delivery Days
1	Manoj Ramanujam	110029,Ansari Nagar	1800000	685

Special terms and conditions-Version:1 effective from 06-10-2025 for category Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile and Single Use) Conforming to IS/ISO 10555 (Part 5)

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. **Turnover**

   Bidder Turn Over Criteria: The minimum average annual financial turnover of the bidder during the last three years, ending on 31st March of the previous financial year, should be as indicated in the bid document. Documentary evidence in the form of certified Audited Balance Sheets of relevant periods or a certificate from the Chartered Accountant / Cost Accountant indicating the turnover details for the relevant period shall be uploaded with the bid. In case the date of constitution / incorporation of the bidder is less than 3 year old, the average turnover in respect of the completed financial years after the date of constitution shall be taken into account for this criteria.

5. **Turnover**

   OEM Turn Over Criteria: The minimum average annual financial turnover of the OEM of the offered product during the last three years, ending on 31st March of the previous financial year, should be as indicated in the bid document. Documentary evidence in the form of certified Audited Balance Sheets of relevant periods or a certificate from the Chartered Accountant / Cost Accountant indicating the turnover details for the relevant period shall be uploaded with the bid. In case the date of constitution / incorporation of the OEM is less than 3 year old, the average turnover in respect of the completed financial years after the date of constitution shall be taken into account for this criteria. In case of bunch bids, the OEM of CATEGORY RELATED TO primary product having highest bid value should meet this criterion.

6. **OEM**

   IMPORTED PRODUCTS: In case of imported products, OEM or Authorized Seller of OEM should have a registered office in India to provide after sales service support in India. The certificate to this effect should be submitted.

7. **Financial Criteria**

   NET WORTH: Net Worth of the OEM should be positive as per the last audited financial statement.

8. **Certificates**

   Bidder's offer is liable to be rejected if they don't upload any of the certificates / documents sought in the Bid document, ATC and Corrigendum if any.

9. **Certificates**

   ISO 9001: The bidder or the OEM of the offered products must have ISO 9001 certification.

10. **Certificates**

    Material Test Certificate Should Be Sent Along with The Supply. The Material Will Be Checked by Buyer's Lab & the Results of the Lab will be the Sole Criteria for Acceptance of the Item.

11. **Certificates**

    The bidder is required to upload, along with the bid, all relevant certificates such as BIS licence, type test certificate, approval certificates and other certificates as prescribed in the Product Specification given in the bid document.

12. **Certificates**

    To be eligible for award of contract, Bidder / OEM must possess following Certificates / Test Reports on the date of bid opening (to be uploaded with bid):

    AS MENTIONED UNDER BUYER ADDED BID SPECIFIC ATC (Click the link to see (in blue color))
    .

13. **Generic**

    Bidder financial standing: The bidder should not be under liquidation, court receivership or similar proceedings, should not be bankrupt. Bidder to upload undertaking to this effect with bid.

14. **Generic**

    Bidders shall quote only those products (Part of Service delivery) in the bid which are not obsolete in the market and has at least 5 years residual market life i.e. the offered product shall not be declared end-of-life by the OEM before this period.

15. **Generic**

    Data Sheet of the product(s) offered in the bid, are to be uploaded along with the bid documents. Buyers can match and verify the Data Sheet with the product specifications offered. In case of any unexplained mismatch of technical parameters, the bid is liable for rejection.

16. **Generic**

    Experience Criteria: The Bidder or its OEM {themselves or through reseller(s)} should have regularly, manufactured and supplied same or similar Category Products to any Central / State Govt Organization / PSU for 3 years before the bid opening date. Copies of relevant contracts to be submitted along with bid in support of having supplied some quantity during each of the year. In case of bunch bids, the primary product having highest value should meet this criterion.

17. **Generic**

    Manufacturer Authorization: Wherever Authorised Distributors/service providers are submitting the bid, Authorisation Form /Certificate with OEM/Original Service Provider details such as name, designation, address, e-mail Id and Phone No. required to be furnished along with the bid

18. **Generic**

    Products supplied shall be nontoxic and harmless to health. In the case of toxic materials, Material Safety Data Sheet may be furnished along with the material.

19. **Generic**

    Without prejudice to Buyer's right to price adjustment by way of discount or any other right or remedy available to Buyer, Buyer may terminate the Contract or any part thereof by a written notice to the Seller, if:
    i) The Seller fails to comply with any material term of the Contract.
    ii) The Seller informs Buyer of its inability to deliver the Material(s) or any part thereof within the stipulated Delivery Period or such inability otherwise becomes apparent.

iii) The Seller fails to deliver the Material(s) or any part thereof within the stipulated Delivery Period and/or to replace/rectify any rejected or defective Material(s) promptly.
iv) The Seller becomes bankrupt or goes into liquidation.
v) The Seller makes a general assignment for the benefit of creditors.
vi) A receiver is appointed for any substantial property owned by the Seller.
vii) The Seller has misrepresented to Buyer, acting on which misrepresentation Buyer has placed the Purchase Order on the Seller.

20. **Generic**

1. The Seller shall not assign the Contract in whole or part without obtaining the prior written consent of buyer.
2. The Seller shall not sub-contract the Contract in whole or part to any entity without obtaining the prior written consent of buyer.
3. The Seller shall, notwithstanding the consent and assignment/sub-contract, remain jointly and severally liable and responsible to buyer together with the assignee/ sub-contractor, for and in respect of the due performance of the Contract and the Sellers obligations there under.

21. **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 (Increased quantity ÷ Original quantity) × 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.

22. **Buyer Added Bid Specific ATC**

Buyer Added text based ATC clauses

A) Buyer Required Specifications:

Category	Specification	Bid Requirement (Allowed Values)
PACKING	Type of Packing	All Components Packed Together in a Transparent Pack
CERTIFICATIONS & REGULATORY COMPLIANCE	Compliance to Medical Device Rules (MDR) 2017 as amended till date & Medical Device Manufacturing Licence under MDR-2017.	Yes
	Valid Wholesale Drug License (Form 20B/21B or equivalent) (if bidder is OEM authorized distributor)	Yes
	Manufacturing unit certification (Quality Management System as per ISO 13485)	Yes

(Compliance to equivalent or higher standards shall be acceptable)	Availability of Test Report for each supplied batch/product as per Medical Device Rule (MDR) 2017 as amended till date	Yes
	Submission of all necessary certifications, licenses and test reports to the buyer at the time of bid submission or along with supplies as per buyer requirement	Yes
SHELF LIFE	Minimum shelf life of the product at the time of delivery to the consignee	3/4 th of Total Shelf Life
ADVANCE SAMPLE	Agree to provide advance sample of the product for buyer's approval before commencement of supply in case of bidding	Yes

B) Additional Requirements

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1) Sterile, non-pyrogenic, and single-use peripheral intravenous cannula intended for venous access in adult patients for infusion of fluids, blood products, and administration of medications etc.,
2) The IV Cannula shall conform to any one or more of the following standards/regulations, as applicable:
    a) IS/ISO 10555-1 – Intravascular catheters: General requirements
    b) IS/ISO 10555-5 – Over-the-needle peripheral catheters (IV cannula)
    c) Material shall comply with ISO 10993 for biological safety
3) All components should be sterile, only.
4) **Required Size Ranges** (Each size should be clearly color-coded and labelled): 14G, 16G, 18G, 20G and 22G
5) Sterilized by ETO / Gamma radiation or equivalent validated method and should be as per ISO or equivalent methods
6) Individually packed in peel-open blister pack or equivalent
7) Each unit pack shall mention:
    i. Product name
    ii. Gauge size
    iii. Sterilization method
    iv. Batch / Lot number
    v. Date of manufacture
    vi. Expiry date

vii. Manufacturer’s name and address

viii. Any other instructions

8) **Catheter:**

a) Made of medical-grade biocompatible polymer PTFE (Polytetrafluoroethylene) or equivalent biocompatible material suitable for intravascular use

b) Chemically inert and non-toxic

c) Resistant to kinking and deformation during insertion and dwell time

d) Smooth surface finish to minimize vessel trauma and thrombophlebitis etc.,

e) Low friction coefficient for easy insertion

f) Thin-walled construction to provide optimal flow rate for the given gauge size

g) Maintains dimensional stability under normal infusion pressure

h) Sterility and labeling as per Indian Pharmacopoeia (IP)

9) **Introducer Needle:**

a) Made of medical-grade stainless steel or equivalent biocompatible material s uitable for intravascular use.

b) Precision-ground bevelled tip for smooth venipuncture.

c) The needle shall be coaxially aligned with the PTFE catheter to ensure smooth catheter advancement without shearing or damage.

d) The needle shall possess adequate tensile strength and rigidity to prevent be nding, breakage, or detachment during insertion and withdrawal

e) The needle shall be securely bonded to the hub to prevent accidental separation during use.

f) The needle shall allow unobstructed blood flashback into the flashback chamb er upon successful venous entry.

g) The needle shall comply ISO 7864 or equivalent standards for Sterile hypoder mic needles for single use

10) **Flashback Chamber:**

a) The IV cannula shall be provided with an integrated transparent flashback ch amber to enable clear and immediate visualization of blood return upon successf ul venous access.

b) The flashback chamber shall be made of medical-grade, clear polymer that d oes not discolor or become opaque during the shelf life of the product.

c) The design shall ensure unobstructed and rapid blood flashback without air e ntrapment or leakage.

d) The flashback chamber shall be securely bonded to the cannula assembly to prevent detachment during use.

e) The chamber shall be designed to prevent blood spillage during needle withdr awal.

f) The chamber shall allow safe handling and disposal of the cannula after use.

11) **Hub & Injection Port:**

a) The IV cannula shall be provided with a medical-grade polymer hub that is se curely bonded to the catheter to ensure structural integrity during insertion and u se.

b) The hub shall be color-coded according to gauge size in line with internationa lly accepted norms for easy identification.

c) The hub design shall permit secure connection with standard Luer lock fittings of IV infusion sets and accessories.

d) The cannula shall be provided with an integrated injection port fitted with a sel f-sealing, latex-free valve to allow intermittent drug administration without needl e puncture.

e) The injection port shall be leak-proof under normal infusion and injection pres sures and shall prevent blood backflow when not in use.

f) The hub and injection port assembly shall allow one-handed operation, where a pplicable, to facilitate ease of use in clinical settings.

g) The hub and injection port shall be transparent or semi-transparent, permittin g visual confirmation of fluid flow and blood presence.

12) **Stabilization / Wings:**

a) The IV cannula shall be provided with integrated stabilization wings (butterfly type) or an equivalent stabilization platform to facilitate secure fixation to the pat ient's skin.

b) The stabilization wings/platform shall be made of medical-grade, flexible poly mer to ensure patient comfort and reduce the risk of dislodgement.

c) The design shall allow proper alignment of the cannula during insertion and s hall help minimize catheter movement during dwell time.

d) The wings/platform shall be adequately sized and shaped to permit effective t aping or dressing without exerting undue pressure on the insertion site.

e) The stabilization feature shall not interfere with:

i. Catheter advancement

ii. Needle withdrawal

iii. Connection of IV sets or accessories

f) The stabilization wings/platform shall be securely bonded to the hub/catheter assembly and shall not detach during normal use.

13) Estimated bid price will be as per Last purchased prices of Surgical Store-1, Hos pital Store.

14) **Manufactures of the quoted product should submit a duly signed undert aking or self-declaration as part of their bid, explicitly confirming full co mpliance with all the technical specifications outlined above. Submissio n of this declaration is a mandatory prerequisite for bid consideration. F ailure to provide the undertaking will result in outright disqualification of the bid, and no further correspondence or clarification in this regard shall be entertained.**

INFORMATION TO BIDDERS:

1. It is hereby informed that all prospective bidders are required to carefully review the **Buyer Added Bid S pecific Additional Terms and Conditions (ATC),** which is attached below and kindly click the link to ch eck.
2. The ATC document outlines buyer required specifications and detailed terms & conditions which must be a

dhered by all bidders during the bid submission process. Bidders must ensure that their proposals are fully compliant with the specifications and conditions mention ed in the ATC.

3. Non-compliance with any of the stipulated terms may result in disqualification or rejection of the bid.
4. For any further clarification regarding the terms, conditions, or specifications mentioned in the ATC, contac t the below address:

   SURGICAL STORE-I,
   HOSPITAL STORES,
   AIIMS, NEW DELHI
   email: manojr@aiims.gov.in

5. This notice is issued in the interest of maintaining transparency and ensuring that all bidders have equal a ccess to the necessary information.

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23. **Buyer Added Bid Specific ATC**

    Buyer uploaded ATC document Click here to view the file.

अस्वीकरण/Disclaimer

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.
15. Buyer added ATC Clauses which are in contravention of clauses defined by buyer in system generated bid template as indicated above in the Bid Details section, EMD Detail, ePBG Detail and MII and MSE Purchase Preference sections of the bid, unless otherwise allowed by GeM GTC.
16. 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 catalogs 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 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.

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.

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 कार्रवाई 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---