

**बिड दस्तावेज़ / Bid Document**

बिड विवरण/Bid Details	
बिड बंद होने की तारीख/समय /Bid End Date/Time	21-02-2026 13:00:00
बिड खुलने की तारीख/समय /Bid Opening Date/Time	21-02-2026 13:30:00
बिड पेशकश वैधता (बंद होने की तारीख से)/Bid Offer Validity (From End Date)	45 (Days)
मंत्रालय/राज्य का नाम/Ministry/State Name	Ministry Of Labour And Employment
विभाग का नाम/Department Name	Employees State Insurance Corporation
संगठन का नाम/Organisation Name	Employees State Insurance Corporation
कार्यालय का नाम/Office Name	Esic Hospital Rudrapur
कुल मात्रा/Total Quantity	2387
वस्तु श्रेणी /Item Category	Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile and Single Use) Conforming to IS/ISO 10555 (Part 5) (Q2) , Single Use Sterile Hypodermic Syringes for Human Use Conforming to IS 10258 (Part 1) (Q2)
बिडर का न्यूनतम औसत वार्षिक टर्नओवर (3 वर्षों का) /Minimum Average Annual Turnover of the bidder (For 3 Years)	3 Lakh (s)
मूल उपकरण निर्माता का औसत टर्नओवर (गत 3 वर्षों का)/OEM Average Turnover (Last 3 Years)	23 Lakh (s)
उन्हीं/समान सेवा के लिए अपेक्षित विगत अनुभव के वर्ष/Years of Past Experience Required for same/similar service	2 Year (s)
वर्षों के अनुभव एवं टर्नओवर से एमएसई को छूट प्राप्त है / MSE Relaxation for Years Of Experience and Turnover	Yes   Partial   Experience - 1 year (s)   Turn over value - 3 (in lakhs)
स्टार्टअप के लिए अनुभव के वर्षों और टर्नओवर से छूट प्रदान की गई है / Startup Relaxation for Years Of Experience and Turnover	Yes   Partial   Experience - 1 year (s)   Turn over value - 3 (in lakhs)
विक्रेता से मांगे गए दस्तावेज़/Document required from seller	Experience Criteria,Past Performance,Bidder Turnover,Certificate (Requested in ATC),OEM Authorization Certificate,OEM Annual Turnover *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

**बिड विवरण/Bid Details**

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
ऑटो एक्सटेंशन अधिकतम कितनी बार किया जाना है। / Number of Auto Extension count	3
विगत प्रदर्शन /Past Performance	50 %
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	Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile and Single Use) Conforming to IS/ISO 10555 (Part 5)
तकनीकी मूल्यांकन के दौरान तकनीकी स्पष्टीकरण हेतु अनुमत समय /Time allowed for Technical Clarifications during technical evaluation	3 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	Total value wise evaluation
मध्यस्थता खंड/Arbitration Clause	No
सुलह खंड/Mediation Clause	No

**ईएमडी विवरण/EMD Detail**

एडवाइजरी बैंक/Advisory Bank	State Bank of India
ईएमडी राशि/EMD Amount	28000

**ईपीबीजी विवरण /ePBG Detail**

एडवाइजरी बैंक/Advisory Bank	State Bank of India
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ईपीबीजी प्रतिशत (%) / ePBG Percentage (%)	5.00
ईपीबीजी की आवश्यक अवधि (माह) / Duration of ePBG required (Months).	62

(a). जेम की शर्तों के अनुसार ईएमडी छूट के इच्छुक बिडर को संबंधित कटेगरी के लिए बिड के साथ वैध समर्थित दस्तावेज़ 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). 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 :**

MEDICAL SUPERINTENDENT

ESIC Hospital Rudrapur, RESERVE POLICE LINE ROAD, BEHIND MANOJ SARKAR STADIUM, UDHAM SINGH NAGAR, RUDRAPUR, UTHARAKHAND - 263153

(Esic Hospital Rudrapur)

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
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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. 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 above 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.
6. Experience Criteria: In respect of the filter applied for experience criteria, the Bidder or its OEM of the product offered in the bid {themselves or through reseller(s)} should have regularly, manufactured and supplied same or similar Category Products to any Central / State Govt Organization / PSU for number of Financial years as indicated above in the bid document before the bid opening date. Copies of relevant contracts and delivery acceptance certificates like CRAC to be submitted along with bid in support of having supplied some quantity during each of the Financial year. In case of bunch bids, the category of primary product having highest value should meet this criterion.
7. 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.
8. 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.
9. 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.

10. Past Performance: The Bidder or its OEM {themselves or through re-seller(s)} should have supplied same or similar Category Products for 50% of bid quantity, in at least one of the last three Financial years before the bid opening date to any Central / State Govt Organization / PSU. Copies of relevant contracts (proving supply of cumulative order quantity in any one financial year) to be submitted along with bid in support of quantity supplied in the relevant Financial year. In case of bunch bids, the category related to primary product having highest bid value should meet this criterion.

11. 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) ( 2 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 कैटेगरी विशिष्टि के अनुसार / As per GeM Category Specification](#)

<b>विवरण/Specification</b>	<b>विशिष्टि का नाम /Specification Name</b>	<b>बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)</b>
<b>PRODUCT INFORMATION</b>	With Wings	Yes
	Injection Port integral with the catheter hub	Yes
	Protection against accidental needle stick injuries	Yes
	<b>Number of Catheter Lumen</b>	Single Lumen
	<b>Nominal outside diameter (mm)</b>	1.6 mm
	<b>Color coding (as per Table 1 of IS/ISO 10555-5)</b>	Medium Grey
<b>PACKAGING</b>	<b>Number of pieces in a Pack</b>	100

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
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	<b>Compliance to Drugs and Cosmetic Act 1940 and Medical Device Rules (MDR) 2017 as amended till date</b>	Yes
	<b>Availability of valid Medical Device license for the product issued from the competent authority</b>	Yes

**Additional Specification Parameters - Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile And Single Use) Conforming To IS/ISO 10555 (Part 5) ( 2 packet )**

Specification Parameter Name	Bid Requirement (Allowed Values)
warranty	5 years

\* 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	मात्रा /Quantity	डिलीवरी के दिन/Delivery Days
1	Kamal Rai Singh	263153,ESIC Hospital Near police lines, Opp Rudrapur Stadium, Rudrapur,Udham Singh Nagar,Uttarakhand	2	15

**Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile And Single Use) Conforming To IS/ISO 10555 (Part 5) ( 23 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	With Wings	Yes
	Injection Port integral with the catheter hub	Yes

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
	Protection against accidental needle stick injuries	Yes
	<b>Number of Catheter Lumen</b>	Single Lumen
	<b>Nominal outside diameter (mm)</b>	1.3 mm
	<b>Color coding (as per Table 1 of IS/ISO 10555-5)</b>	Deep Green
PACKAGING	<b>Number of pieces in a Pack</b>	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	<b>Compliance to Drugs and Cosmetic Act 1940 and Medical Device Rules (MDR) 2017 as amended till date</b>	Yes
	<b>Availability of valid Medical Device license for the product issued from the competent authority</b>	Yes

**Additional Specification Parameters - Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile And Single Use) Conforming To IS/ISO 10555 (Part 5) ( 23 packet )**

Specification Parameter Name	Bid Requirement (Allowed Values)
warranty	5 years

\* 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	मात्रा /Quantity	डिलीवरी के दिन/Delivery Days
1	Kamal Rai Singh	263153,ESIC Hospital Near police lines, Opp Rudrapur Stadium, Rudrapur,Udham Singh Nagar,Uttarakhand	23	15

**Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile And Single Use) Conforming To IS/ISO 10555 (Part 5) ( 105 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	With Wings	Yes
	Injection Port integral with the catheter hub	Yes
	Protection against accidental needle stick injuries	Yes
	<b>Number of Catheter Lumen</b>	Single Lumen
	<b>Nominal outside diameter (mm)</b>	1.1 mm
	<b>Color coding (as per Table 1 of IS/ISO 10555-5)</b>	Pink
PACKAGING	<b>Number of pieces in a Pack</b>	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	<b>Compliance to Drugs and Cosmetic Act 1940 and Medical Device Rules (MDR) 2017 as amended till date</b>	Yes
	<b>Availability of valid Medical Device license for the product issued from the competent authority</b>	Yes

**Additional Specification Parameters - Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile And Single Use) Conforming To IS/ISO 10555 (Part 5) ( 105 packet )**

Specification Parameter Name	Bid Requirement (Allowed Values)
warranty	5 years

\* 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	मात्रा /Quantity	डिलीवरी के दिन/Delivery Days
1	Kamal Rai Singh	263153,ESIC Hospital Near police lines, Opp Rudrapur Stadium, Rudrapur,Udham Singh Nagar,Uttarakhand	105	15

**Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile And Single Use) Conforming To IS/ISO 10555 (Part 5) ( 110 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	With Wings	Yes
	Injection Port integral with the catheter hub	Yes
	Protection against accidental needle stick injuries	Yes
	<b>Number of Catheter Lumen</b>	Single Lumen
	<b>Nominal outside diameter (mm)</b>	0.8 mm
	<b>Color coding (as per Table 1 of IS/ISO 10555-5)</b>	Deep Blue
PACKAGING	<b>Number of pieces in a Pack</b>	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	<b>Compliance to Drugs and Cosmetic Act 1940 and Medical Device Rules (MDR) 2017 as amended till date</b>	Yes
	<b>Availability of valid Medical Device license for the product issued from the competent authority</b>	Yes

**Additional Specification Parameters - Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile And Single Use) Conforming To IS/ISO 10555 (Part 5) ( 110 packet )**

Specification Parameter Name	Bid Requirement (Allowed Values)
warranty	5 years

\* 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	Kamal Rai Singh	263153,ESIC Hospital Near police lines, Opp Rudrapur Stadium, Rudrapur,Udham Singh Nagar,Uttarakhand	110	15

**Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile And Single Use) Conforming To IS/ISO 10555 (Part 5) ( 50 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	With Wings	Yes
	Injection Port integral with the catheter hub	Yes
	Protection against accidental needle stick injuries	Yes
	<b>Number of Catheter Lumen</b>	Single Lumen
	<b>Nominal outside diameter (mm)</b>	0.7 mm
	<b>Color coding (as per Table 1 of IS/ISO 10555-5)</b>	Yellow
PACKAGING	<b>Number of pieces in a Pack</b>	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	<b>Compliance to Drugs and Cosmetic Act 1940 and Medical Device Rules (MDR) 2017 as amended till date</b>	Yes
	<b>Availability of valid Medical Device license for the product issued from the competent authority</b>	Yes

**Additional Specification Parameters - Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile And Single Use) Conforming To IS/ISO 10555 (Part 5) ( 50 packet )**

Specification Parameter Name	Bid Requirement (Allowed Values)
warranty	5 years

\* 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	Kamal Rai Singh	263153,ESIC Hospital Near police lines, Opp Rudrapur Stadium, Rudrapur,Udham Singh Nagar,Uttarakhand	50	15

**Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile And Single Use) Conforming To IS/ISO 10555 (Part 5) ( 32 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 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	With Wings	Yes
	Injection Port integral with the catheter hub	Yes
	Protection against accidental needle stick injuries	Yes
	<b>Number of Catheter Lumen</b>	Single Lumen
	<b>Nominal outside diameter (mm)</b>	0.6 mm
	<b>Color coding (as per Table 1 of IS/ISO 10555-5)</b>	Violet
PACKAGING	<b>Number of pieces in a Pack</b>	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	<b>Compliance to Drugs and Cosmetic Act 1940 and Medical Device Rules (MDR) 2017 as amended till date</b>	Yes
	<b>Availability of valid Medical Device license for the product issued from the competent authority</b>	Yes

**Additional Specification Parameters - Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile And Single Use) Conforming To IS/ISO 10555 (Part 5) ( 32 packet )**

Specification Parameter Name	Bid Requirement (Allowed Values)
warranty	5 years

\* 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	मात्रा /Quantity	डिलीवरी के दिन/Delivery Days
1	Kamal Rai Singh	263153,ESIC Hospital Near police lines, Opp Rudrapur Stadium, Rudrapur,Udham Singh Nagar,Uttarakhand	32	15

**Single Use Sterile Hypodermic Syringes For Human Use Conforming To IS 10258 (Part 1) ( 700 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 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	<b>Needle included</b>	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
	<b>Nominal Capacity of Syringe (ml)</b>	< 2
PACKAGING	<b>Number of pieces in a Pack</b>	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	<b>Compliance to Drugs and Cosmetic Act 1940 and Medical Device Rules (MDR) 2017 as amended till date</b>	Yes

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
	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) ( 700 packet )**

Specification Parameter Name	Bid Requirement (Allowed Values)
warranty	5 years
needle size	24G

\* 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	मात्रा /Quantity	डिलीवरी के दिन/Delivery Days
1	Kamal Rai Singh	263153,ESIC Hospital Near police lines, Opp Rudrapur Stadium, Rudrapur,Udham Singh Nagar,Uttarakhand	700	15

**Single Use Sterile Hypodermic Syringes For Human Use Conforming To IS 10258 (Part 1) ( 800 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 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	<b>Needle included</b>	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
	<b>Nominal Capacity of Syringe (ml)</b>	>2 - 5

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
PACKAGING	Number of pieces in a Pack	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 - Single Use Sterile Hypodermic Syringes For Human Use Conforming To IS 10258 (Part 1) ( 800 packet )**

Specification Parameter Name	Bid Requirement (Allowed Values)
WARRANTY	5 YEARS
NEEDLE SIZE	24G

\* 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	Kamal Rai Singh	263153,ESIC Hospital Near police lines, Opp Rudrapur Stadium, Rudrapur,Udham Singh Nagar,Uttarakhand	800	15

**Single Use Sterile Hypodermic Syringes For Human Use Conforming To IS 10258 (Part 1) ( 500 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	<b>Needle included</b>	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
	<b>Nominal Capacity of Syringe (ml)</b>	>5 - 10
PACKAGING	<b>Number of pieces in a Pack</b>	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	<b>Compliance to Drugs and Cosmetic Act 1940 and Medical Device Rules (MDR) 2017 as amended till date</b>	Yes
	<b>Availability of valid Medical Device license for the product issued from the competent authority</b>	Yes

**Additional Specification Parameters - Single Use Sterile Hypodermic Syringes For Human Use Conforming To IS 10258 (Part 1) ( 500 packet )**

Specification Parameter Name	Bid Requirement (Allowed Values)
WARRANTY	5 YEARS
NEEDLE SIZE	24G

\* 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	मात्रा /Quantity	डिलीवरी के दिन/Delivery Days
1	Kamal Rai Singh	263153,ESIC Hospital Near police lines, Opp Rudrapur Stadium, Rudrapur,Udham Singh Nagar,Uttarakhand	500	15

**Single Use Sterile Hypodermic Syringes For Human Use Conforming To IS 10258 (Part 1) ( 60 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	<b>Needle included</b>	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
	<b>Nominal Capacity of Syringe (ml)</b>	>10 - 20
PACKAGING	<b>Number of pieces in a Pack</b>	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	<b>Compliance to Drugs and Cosmetic Act 1940 and Medical Device Rules (MDR) 2017 as amended till date</b>	Yes
	<b>Availability of valid Medical Device license for the product issued from the competent authority</b>	Yes

**Additional Specification Parameters - Single Use Sterile Hypodermic Syringes For Human Use Conforming To IS 10258 (Part 1) ( 60 packet )**

Specification Parameter Name	Bid Requirement (Allowed Values)
WARRANTY	5 YEARS

\* 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	Kamal Rai Singh	263153,ESIC Hospital Near police lines, Opp Rudrapur Stadium, Rudrapur,Udham Singh Nagar,Uttarakhand	60	15

## Single Use Sterile Hypodermic Syringes For Human Use Conforming To IS 10258 (Part 1) ( 5 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	<b>Needle included</b>	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
	<b>Nominal Capacity of Syringe (ml)</b>	? 50
PACKAGING	<b>Number of pieces in a Pack</b>	50
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	<b>Compliance to Drugs and Cosmetic Act 1940 and Medical Device Rules (MDR) 2017 as amended till date</b>	Yes
	<b>Availability of valid Medical Device license for the product issued from the competent authority</b>	Yes

### Additional Specification Parameters - Single Use Sterile Hypodermic Syringes For Human Use Conforming To IS 10258 (Part 1) ( 5 packet )

Specification Parameter Name	Bid Requirement (Allowed Values)
WARRANTY	5 YEARS

\* 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	मात्रा /Quantity	डिलीवरी के दिन/Delivery Days

क्र.सं./S.No.	प्रेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer	पता/Address	मात्रा /Quantity	डिलीवरी के दिन/Delivery Days
1	Kamal Rai Singh	263153,ESIC Hospital Near police lines, Opp Rudrapur Stadium, Rudrapur,Udham Singh Nagar,Uttarakhand	5	15

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

**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](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. **Generic**

OPTION CLAUSE: The Purchaser reserves the right to increase or decrease the quantity to be ordered up to 25 percent of bid quantity at the time of placement of contract. The purchaser also reserves the right to increase the ordered quantity up to 25% 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.

### 2. **Sample Clause**

After award of contract - Successful Bidder shall have to get advance sample approved from buyer before bulk manufacturing / starting bulk supplies. Successful Bidder shall submit

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samples for Buyer's approval, within 7 days of award of contract. Buyer shall, as per contract specifications framework, either approve the advance sample or will provide complete list of modification required in the sample within 15 days of receipt of advance sample. Seller shall be required to ensure supply as per approved sample with modifications as communicated by Buyer. If there is delay from buyer side in approval of advance sample - the delivery period shall be refixed without LD for the period of delay in sample approval. In case, the sample is found to have major deviations / not conforming to the Contract specifications, the buyer at its discretion may call for fresh samples for approval before allowing bulk supplies or may terminate the contract after notifying the deviations to the seller.

Unless otherwise provided in the contract, all samples required for test shall be supplied by the contractor free of cost. Where under the contract, the contractor is required to submit an advance sample, any expenses incurred by the contractor on or in connection with the production of stores in bulk, before the sample has been approved unconditionally shall be borne by the Seller and he shall not claim any compensation in the event of such sample being found unacceptable by the Buyer / Consignee.

### 3. **Generic**

Actual delivery (and Installation & Commissioning (if covered in scope of supply)) is to be done at following address

ESIC HOSPITAL RUDRAPUR  
RESERVE POLICE LINE ROAD  
BEHIND MANOJ SARKAR STADIUM  
UDHAM SINGH NAGAR  
RUDRAPUR, UTHARAKHAND -263153

#### 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. **Forms of EMD and PBG**

Bidders can also submit the EMD with Account Payee Demand Draft in favour of

ESIC FUND ACCOUNT NUMBER 1  
payable at  
ESIC RUDRAPUR

Bidder has to upload scanned copy / proof of the DD along with bid and has to ensure delivery of hardcopy to the Buyer within 5 days of Bid End date / Bid Opening date.

#### 6. **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.

#### 7. **Generic**

Bidders are advised to check applicable GST on their own before quoting. Buyer will not take any responsibility in this regards. GST reimbursement will be as per actuals or as per applicable rates (whichever is lower), subject to the maximum of quoted GST %.

#### 8. **Buyer Added Bid Specific ATC**

Buyer Added text based ATC clauses

EXPIRY SHOULD BE 5 YEARS; ONE SAMPLE OF EACH ITEMS SHOULD BE PROVIDED WITH IN ONE WEEK FOR TECHNICAL EVALUATION AFTER THE BID OPENING DATE.

WARRANTY WHEREVER MENTIONED SHOULD BE CONSIDERED AS EXPIRY.

#### 9. **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 3 years residual market life i.e. the offered product shall not be declared end-of-life by the OEM before this period.

#### 10. **Rate Contract**

Performance Security: Against all Purchase Orders placed by any buyer against the Rate Contract Catalogue, the seller would be required to furnish a performance security of 3 % of total contract value of the relevant Purchase Order as per GeM GTC.

11. **Rate Contract**

**Packing Material** The items will be supplied by the successful bidder in its original packing material and the packing material will not be returned. Weight of packing material will not be included in quantity supplied and only net weight of the items will be counted. The packing should be of standardized weights of appropriate size. Item will not be accepted in non-standardized weights. Packing Material. The items will be supplied by the successful bidder in its original packing material and the packing material will not be returned. Weight of packing material will not be included in quantity supplied and only net weight of the items will be counted. The packing should be of standardized weights of appropriate size. Item will not be accepted in non-standardized weights.

12. **Rate Contract**

**Demurrage charges** In case the rejected items are not lifted by the firm within 48 hrs, the demurrage charges at the rate of 0.5% of total contract value will be charged per day. In case the items are not lifted within a month, the same will be destroyed by the station board of officers and no claim will be admitted. Demurrage charges. In case the rejected items are not lifted by the firm within 48 hrs, the demurrage charges at the rate of 0.5% of total contract value will be charged per day. In case the items are not lifted within a month, the same will be destroyed by the station board of officers and no claim will be admitted.

13. **Rate Contract**

Tax and Duties Toll Tax/ Entry Tax/ Octroi Duty & Local Taxes. No separate payment would be made by the Buyer for Toll Taxes, Entry Taxes, Octroi duty and local Taxes, if any. The Seller should cater for these Taxes/ duties as part of Basic Rate quoted in the Bid.

14. **Rate Contract**

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.

15. **Rate Contract**

Bidder's offer is liable to be rejected if they dont upload any of the certificates / documents sought in the Bid document, ATC and Corrigendum if any.

16. **Rate Contract**

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.

17. **Rate Contract**

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.

18. **Rate Contract**

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

19. **Rate Contract**

Bidders are advised to check applicable GST on their own before quoting. Buyer will not take any responsibility in this regards. GST reimbursement will be as per actuals or as per applicable rates (whichever is lower), subject to the maximum of quoted GST %.

## 20. **Warranty**

Warranty period of the supplied products shall be 5 years from the date of final acceptance of goods or after completion of installation, commissioning & testing of goods (if included in the scope of supply), at consignee location. OEM Warranty certificates must be submitted by Successful Bidder at the time of delivery of Goods. The seller should guarantee the rectification of goods in case of any break down during the guarantee period. Seller should have well established Installation, Commissioning, Training, Troubleshooting and Maintenance Service group in INDIA for attending the after sales service. Details of Service Centres near consignee destinations are to be uploaded along with the bid.

## 21. **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.

## 22. **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 2 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.

## 23. **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.

## 24. **Certificates**

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

## 25. **Purchase Preference (Centre)**

Purchase preference to Micro and Small Enterprises (MSEs): Purchase preference will be given to MSEs 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 the Purchase preference, the bidder must be the manufacturer of the offered product in case of bid for supply of goods. Traders are excluded from the purview of Public Procurement Policy for Micro and Small Enterprises. 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. If L-1 is not an MSE and MSE Seller (s) has/have quoted price within L-1+ 15% of margin of purchase preference /price band defined in relevant policy, such Seller shall be given opportunity to match L-1 price and contract will be awarded for percentage of 15% of total value.

## 26. **Past Project Experience**

**Proof for Past Experience and Project Experience clause:** For fulfilling the experience criteria any one of the following documents may be considered as valid proof for meeting the experience criteria:a. Contract copy along with Invoice(s) with self-certification by the bidder that service/supplies against the invoices have been executed.b. Execution certificate by client with contract value.c. Any other document in support of contract execution like Third Party Inspection release note, etc.**Proof for Past Experience and Project Experience clause:** For fulfilling the experience criteria any one of the following documents may be considered as valid proof for meeting the experience criteria:a. Contract copy along with Invoice(s) with self-certification by the bidder that service/supplies against the invoices have been executed.b. Execution certificate by client with contract value.c. Any other document in support of contract execution like Third

Party Inspection release note, etc.

**27. Scope of Supply**

Scope of supply (Bid price to include all cost components) : Only supply of Goods

**28. Forms of EMD and PBG**

Successful Bidder can submit the Performance Security in the form of Account Payee Demand Draft also (besides PBG which is allowed as per GeM GTC). DD should be made in favour of

ESIC FUND ACCOUNT NUMBER 1

payable at

ESIC RUDRAPUR

. After award of contract, Successful Bidder can upload scanned copy of the DD in place of PBG and has to ensure delivery of hard copy to the original DD to the Buyer within 15 days of award of contract.

**29. Generic**

Supplier shall ensure that the Invoice is raised in the name of Consignee with GSTIN of Consignee only.

**30. 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.

**31. 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.

**32. 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):

ALL ASKED IN BID

**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---**