बिड संख्या/Bid Number: GEM/2026/B/8116034

दिनांक /Dated: 05-10-2026

दर अनुबंध बिड दस्तावेज़ / Rate Contract Bid Document

This Rate Contract Bid is governed by General Terms and Conditions of Rate Contract

बिड विवरण/Bid Details	
बिड बंद होने की तारीख/समय /Bid End Date/Time	27-10-2026 16:00:00
बिड खुलने की तारीख/समय /Bid Opening Date/Time	28-10-2026 16:00:00
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	Raipur
शिकायत निवारण के संपर्क विवरण/ Contact details of Grievance redressal	HOD Email id :uttam.sharma36@gov.in Buyer Email id: hospitalstoreskc.rc3@gmail.com
वस्तु श्रेणी /Item Category	Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile and Single Use) Conforming to IS/ISO 10555 (Part 5) (Q2)
बिडर का न्यूनतम औसत वार्षिक टर्नओवर (3 वर्षों का) /Minimum Average Annual Turnover of the bidder (For 3 Years)	17 Lakh (s)
मूल उपकरण निर्माता का औसत टर्नओवर (गत 3 वर्षों का)/OEM Average Turnover (Last 3 Years)	143 Lakh (s)
उन्हीं/समान सेवा के लिए अपेक्षित विगत अनुभव के वर्ष/Years of Past Experience Required for same/similar service	3 Year (s)
टर्नओवर पात्रता मानदंड/Turnover eligibility criteria	GeM verified turnover
स्टार्टअप के लिए अनुभव के वर्षों और टर्नओवर से छूट प्रदान की गई है / Startup Relaxation for Years Of Experience and Turnover	Yes | Complete

बिड विवरण/Bid Details	
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,Additional Doc 1 (Requested in ATC),Additional Doc 2 (Requested in ATC),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	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	1
बिड से रिवर्स नीलामी सक्रिय किया/Bid to RA enabled	No
दर अनुबंध और शीर्षक / Rate Contract & Title	Yes ; IV CANNULA 1.7MM, IV CANNULA 1.3MM, IV CANNULA 1.0MM, IV CANNULA 0.8 &0.9M, IV CANNULA 0.7MM, IV CANNULA 0.6MM, IV CANNULA 0.8 &0.9M, IV CANNULA 0.7MM, IV CANNULA 0.6MM, IV CANNULA 2.2MM
दर अनुबंध की वैधता अवधि / Validity Period Of Rate Contract	12 Months
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
अनुमानित बिड मूल्य / Estimated Bid Value	3591000
मूल्यांकन पद्धति/Evaluation Method	Item wise consignee wise evaluation

ईएमडी विवरण/EMD Detail

एडवाईजरी बैंक/Advisory Bank	Bank of India
ईएमडी राशि/EMD Amount	35000

Discount Slab Range

Over-needle Peripheral Catheters (intravascular Catheters - Sterile And Single Use) Conforming To Is/iso 10555 (part 5)

Slab 1	1 - and above Unit

Over-needle Peripheral Catheters (intravascular Catheters - Sterile And Single Use) Conforming To Is/iso 10555 (part 5)

Slab 1	1 - and above Unit

Over-needle Peripheral Catheters (intravascular Catheters - Sterile And Single Use) Conforming To Is/iso 10555 (part 5)

Slab 1	1 - and above Unit

Over-needle Peripheral Catheters (intravascular Catheters - Sterile And Single Use) Conforming To Is/iso 10555 (part 5)

Slab 1	1 - and above Unit

Over-needle Peripheral Catheters (intravascular Catheters - Sterile And Single Use) Conforming To Is/iso 10555 (part 5)

Slab 1	1 - and above Unit

Over-needle Peripheral Catheters (intravascular Catheters - Sterile And Single Use) Conforming To Is/iso 10555 (part 5)

Slab 1	1 - and above Unit

Over-needle Peripheral Catheters (intravascular Catheters - Sterile And Single Use) Conforming To Is/iso 10555 (part 5)

Slab 1	1 - and above Unit

Over-needle Peripheral Catheters (intravascular Catheters - Sterile And Single Use) Conforming To Is/iso 10555 (part 5)

Slab 1	1 - and above Unit

Over-needle Peripheral Catheters (intravascular Catheters - Sterile And Single Use) Conforming To Is/iso 10555 (part 5)

Slab 1	1 - and above Unit

Over-needle Peripheral Catheters (intravascular Catheters - Sterile And Single Use) Conforming To Is/iso 10555 (part 5)

Slab 1	1 - and above Unit

Over-needle Peripheral Catheters (intravascular Catheters - Sterile And Single Use) Conforming To Is/iso 10555 (part 5)

Slab 1	1 - and above Unit

(a). जेम की शर्तों के अनुसार ईएमडी छूट के इच्छुक बिडर को संबंधित केटेगरी के लिए बिड के साथ वैध समर्थित दस्तावेज़ प्रस्तुत करने हैं। 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 should be in favour of Beneficiary, wherever it is applicable.

(c).ईएमडी और संपादन जमानत राशि लाभार्थी के पक्ष में होनी चाहिए। / Earnest Money Deposit (EMD) shall also be accepted by the buyer in the form of a surety bond.

लाभार्थी /Beneficiary :
AIIMS RAIPUR
Stores Office - Hospital, 1st floor, Block-A, New Administration Building, Gate No. 5, AIIMS Raipur (C.G) Pin-492099
(Aiims Raipur)

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

1. 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.
2. 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.
3. The minimum average annual financial turnover of the bidder during the last three financial years, ending on 31st March of the previous financial year, should be as indicated in the bid document. The turnover of the bidder shall be validated by the system based on the financial data available on the GeM portal. The average turnover shall be calculated based on the last three financial years where data is available; in case data is available for only two of the last three financial years, the average shall be computed based on those two years. In case the date of constitution / incorporation of the bidder is less than three years old, the average turnover in respect of the completed financial years after the date of constitution / incorporation shall be taken into account for this criterion. No separate documentary evidence is required to be uploaded by the bidder for this criterion, as the system shall derive and validate the turnover figures from the data available on the GeM portal.
4. 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.
5. 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.
6. 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.

Pre Bid Detail(s)

मूल्य भिन्नता खंड दस्तावेज़/Pre-Bid Date and Time	प्री-बिड स्थान/Pre-Bid Venue
22-10-2026 15:00:00	Stores Office - Hospital, 1st floor, Block-A, New Administration Building, Gate No. 5, AIIMS Raipur (C.G) Pin-492099

मूल्यांकन विधि(मदवार मूल्यांकन विधि) / **Evaluation Method & Consignee Location** (Item Wise Consignee Wise Evaluation Method)

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

मूल्यांकन अनुसूचियां / **Evaluation Schedules**	वस्तु/श्रेणी / **Item/Category**	**Location of Consignee**
Schedule 1 (RAIPUR)	Over-needle Peripheral Catheters (intravascular Catheters - Sterile And Single Use) Conforming To Is/iso 10555 (part 5)	RAIPUR
Schedule 2 (RAIPUR)	Over-needle Peripheral Catheters (intravascular Catheters - Sterile And Single Use) Conforming To Is/iso 10555 (part 5)	RAIPUR
Schedule 3 (RAIPUR)	Over-needle Peripheral Catheters (intravascular Catheters - Sterile And Single Use) Conforming To Is/iso 10555 (part 5)	RAIPUR

Schedule 4 (RAIPUR)	Over-needle Peripheral Catheters (intravascular Catheters - Sterile And Single Use) Conforming To Is/iso 10555 (part 5)	RAIPUR
Schedule 5 (RAIPUR)	Over-needle Peripheral Catheters (intravascular Catheters - Sterile And Single Use) Conforming To Is/iso 10555 (part 5)	RAIPUR
Schedule 6 (RAIPUR)	Over-needle Peripheral Catheters (intravascular Catheters - Sterile And Single Use) Conforming To Is/iso 10555 (part 5)	RAIPUR
Schedule 7 (RAIPUR)	Over-needle Peripheral Catheters (intravascular Catheters - Sterile And Single Use) Conforming To Is/iso 10555 (part 5)	RAIPUR
Schedule 8 (RAIPUR)	Over-needle Peripheral Catheters (intravascular Catheters - Sterile And Single Use) Conforming To Is/iso 10555 (part 5)	RAIPUR
Schedule 9 (RAIPUR)	Over-needle Peripheral Catheters (intravascular Catheters - Sterile And Single Use) Conforming To Is/iso 10555 (part 5)	RAIPUR
Schedule 10 (RAIPUR)	Over-needle Peripheral Catheters (intravascular Catheters - Sterile And Single Use) Conforming To Is/iso 10555 (part 5)	RAIPUR

Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile And Single Use) Conforming To IS/ISO 10555 (Part 5)

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	Type of Vent Fitting	Fixed fitting, Removable fitting
	With Wings	Yes
	Injection Port integral with the catheter hub	No
	Protection against accidental needle stick injuries	No
	Number of Catheter Lumen	Single Lumen
	Nominal outside diameter (mm)	1.7 mm
	Color coding (as per Table 1 of IS/ISO 10555-5)	Medium Grey
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

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

Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile And Single Use) Conforming To IS/ISO 10555 (Part 5)

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	Type of Vent Fitting	Fixed fitting
	With Wings	Yes
	Injection Port integral with the catheter hub	No
	Protection against accidental needle stick injuries	No
	Number of Catheter Lumen	Single Lumen
	Nominal outside diameter (mm)	1.3 mm
	Color coding (as per Table 1 of IS/ISO 10555-5)	Deep Green
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

Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile And Single Use) Conforming To IS/ISO 10555 (Part 5)

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	Type of Vent Fitting	Fixed fitting, Removable fitting
	With Wings	Yes
	Injection Port integral with the catheter hub	No
	Protection against accidental needle stick injuries	No
	Number of Catheter Lumen	Single Lumen
	Nominal outside diameter (mm)	1.0 mm, 1.1 mm
	Color coding (as per Table 1 of IS/ISO 10555-5)	Pink
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

Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile And Single Use) Conforming To IS/ISO 10555 (Part 5)

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	Type of Vent Fitting	Fixed fitting, Removable fitting
	With Wings	Yes
	Injection Port integral with the catheter hub	No
	Protection against accidental needle stick injuries	No
	Number of Catheter Lumen	Single Lumen
	Nominal outside diameter (mm)	0.8 mm, 0.9 mm
	Color coding (as per Table 1 of IS/ISO 10555-5)	Deep Blue
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

Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile And Single Use) Conforming To IS/ISO 10555 (Part 5)

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	Type of Vent Fitting	Fixed fitting
	With Wings	Yes
	Injection Port integral with the catheter hub	No

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
	Protection against accidental needle stick injuries	No
	Number of Catheter Lumen	Single Lumen
	Nominal outside diameter (mm)	0.7 mm
	Color coding (as per Table 1 of IS/ISO 10555-5)	Yellow
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

Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile And Single Use) Conforming To IS/ISO 10555 (Part 5)

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	Type of Vent Fitting	Fixed fitting
	With Wings	Yes
	Injection Port integral with the catheter hub	No
	Protection against accidental needle stick injuries	No
	Number of Catheter Lumen	Single Lumen
	Nominal outside diameter (mm)	0.6 mm

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
	Color coding (as per Table 1 of IS/ISO 10555-5)	Violet
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

Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile And Single Use) Conforming To IS/ISO 10555 (Part 5)

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	Type of Vent Fitting	Fixed fitting
	With Wings	Yes
	Injection Port integral with the catheter hub	Yes
	Protection against accidental needle stick injuries	No
	Number of Catheter Lumen	Single Lumen
	Nominal outside diameter (mm)	0.8 mm, 0.9 mm
	Color coding (as per Table 1 of IS/ISO 10555-5)	Deep Blue
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

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

Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile And Single Use) Conforming To IS/ISO 10555 (Part 5)

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	Type of Vent Fitting	Fixed fitting
	With Wings	Yes
	Injection Port integral with the catheter hub	Yes
	Protection against accidental needle stick injuries	No
	Number of Catheter Lumen	Single Lumen
	Nominal outside diameter (mm)	0.7 mm
	Color coding (as per Table 1 of IS/ISO 10555-5)	Yellow
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

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

Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile And Single Use) Conforming To IS/ISO 10555 (Part 5)

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	Type of Vent Fitting	Fixed fitting
	With Wings	Yes
	Injection Port integral with the catheter hub	Yes
	Protection against accidental needle stick injuries	No
	Number of Catheter Lumen	Single Lumen
	Nominal outside diameter (mm)	0.6 mm
	Color coding (as per Table 1 of IS/ISO 10555-5)	Violet
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

Over-Needle Peripheral Catheters (Intravascular Catheters - Sterile And Single Use) Conforming To IS/ISO 10555 (Part 5)

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	Type of Vent Fitting	Removable fitting
	With Wings	Yes
	Injection Port integral with the catheter hub	No
	Protection against accidental needle stick injuries	No
	Number of Catheter Lumen	Single Lumen
	Nominal outside diameter (mm)	2.0 mm
	Color coding (as per Table 1 of IS/ISO 10555-5)	Orange
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

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. *A*ll Provisions of Drugs and Cosmetics Act, 1940 and Rules (including Medical Device Rule 2017) made there under and all subsequent amendments till

date will always be applicable. This will include all notifications issued by Central Drugs Standard Control Organization (CDSCO), Ministry of Health & Family Welfare (MoHFW) and Department of Pharmaceuticals (DOP), Ministry of Chemicals & Fertilizers in this regard.

2. The seller’s registration and product upload on GeM is based on their declaration of possessing a valid Medical Device License for the product, certifications, test reports as per MDR (2017), latest amended. However, buyers must check, validate and verify the details at their end for all applicable licenses and certifications e.g., validity and authenticity/genuineness of medical device license (or registration), product certifications, manufacturer’s certifications/ licenses, test reports at the time of supply.
3. In case of authorized resellers/distributors, it will be the legal and regulatory liability of the manufacturer to ensure that their resellers/distributors are operating in compliance with all relevant laws and regulations, and are licensed to sell the manufacturer's products, including verifying the validity and authenticity of license held by them.
4. Any other Terms and Conditions which is not included or at variance with the conditions specified in STC/GTC, may be added by the buyer through Additional Terms and Conditions (ATC) in the bid to ensure items are procured from authentic/validated source with appropriate and applicable quality. The above terms and conditions are in reverse order of precedence i.e. BID ATC shall supersede specific STC which shall supersede General Terms and Conditions (GTC), whenever there are any conflicting provisions.
5. **Packing and Marking:** Should be as per MDR.
6. Presently, the products under this category are not covered under mandatory BIS licensing scheme. Hence, both ISI Marked and non-ISI Marked products may be available in this category. However, if a buyer intends to procure ISI Marked products, buyer may, at his discretion, opt for the same by indicating the requirement in the bid, after taking into account the number of valid licenses issued under the standard. In such cases, buyer shall verify valid BIS License as per the applicable governing standard at the time of evaluation of bid and check the ISI marking on the products at the time of receipt of material before issuing of CRAC.
   The valid BIS license status of sellers may be viewed by buyer from the below link.

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

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

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

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

3. Make in india specific authorisation certificate needs to be enclosed.
4. **Generic**

   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.

5. **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.

6. **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.

7. **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

8. **Scope of Supply**

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

9. **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.

10. **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.

11. **Forms of EMD and PBG**

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

AIIMS Raipur
payable at
AIIMS Raipur
.
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.

12. 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

AIIMS Raipur
payable at
AIIMS Raipur
. 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.

13. Buyer Added Bid Specific ATC

Buyer uploaded ATC document Click here to view the file.

14. 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.

15. 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.

16. 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.

17. 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.

18. 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.

19. 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.

20. 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 %.

21. Buyer Added Bid Specific ATC

Buyer Added text based ATC clauses

STANDARDIZATION OF SPECIFICATIONS Product Specification IV cannula without Injection Port

a. I.V Cannula with wing & without Injection port.

b. Kink resistant, FEP catheter material

c. 4 Radio-opaque contrast lines for X-Ray visualization.

d. Tapered molded Cannula tip.

e. Super sharp triple facetted beveled needle.

f. Transparent flashback chamber allows immediate detection of blood.

g. Sterile, individually packed in blister pack.

STANDARDIZATION OF SPECIFICATIONS Product Specification IV cannula with Injection port

a) I.V Cannula with wing,

b) Kink resistant, FEP catheter material

c) Centrally placed injection port between the wings.

d) Injection port with one way valve.

e) 4 radio-opaque contrast lines for X-Ray visualization.

f) Tapered molded cannula tip.

g) Super sharp triple facetted beveled needle.

h) Transparent flashback chamber allows immediate detection of blood.

i) Sterile, individually packed in blister pack Sample Submission The Participant must submit the minimum 100 pcs samples of th e per quoted products for technical evaluation within a week fro m the last date of submission of bid with mentioning of bid No. And product name.

Additional Requirements

1. The product shall be a sterile, non-pyrogenic, single-use peripheral intravenous cannula intended for venous access in adult patients for infusion of fluids, blood products, and administration of medications.

2. The IV Cannula shall conform to one or more of the following standards/regulations, as applicable: a) IS/ISO 10555-1 – *Intravascular catheters: General r equirements*
   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 shall be sterile only.

4. Required size ranges (each size clearly color-coded and labelled): **14G, 16G, 18G, 20G, 22G, 24G and 26G**.

5. Cannula shall be sterilized by ETO / Gamma radiation or equivalent validated method, in compliance with ISO or equivalent standards.

6. Each unit shall be individually packed in a peel-open blister pack or equivalent.

7. Each unit pack shall clearly mention:
   - Product name
   - Gauge size
   - Sterilization method
   - Batch / Lot number
   - Date of manufacture
   - Expiry date
   - Manufacturer’s name and address
   - Any other relevant instructions

Injection Port:

8. a) The IV cannula shall be provided with a medical-grade polymer hub that is securely bonded to the catheter to ensure structural integrity during insertion and use. b) The hub shall be color-coded accor ding to gauge size in line with internationally accepted norms for easy identification.

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

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

   e) The injection port shall be leak-proof under normal infusion and injection pressures and shall preve nt blood backflow when not in use.

   f) The hub and injection port assembly shall allow one-handed operation, where applicable, to

facilitate ease of use in clinical settings.

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

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अस्वीकरण/Disclaimer

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

1. Publishing Custom / BOQ bids for items for which regular GeM categories are available (unless such Custom / BOQ item is bunched with the major regular product Category Item).
2. Mandating procurement of / from specific Brand / Make / Model / Manufacturer / Dealer except in case of Single Bid / Proprietary Article Certificate (PAC) Buying.
3. Inclusion of disqualification criteria related to suspension of seller / service provider, where such suspension period has already expired.
4. Mandating submission of documents (other than EMD and signed Integrity Pact - which can be submitted within 5 days of bid opening) in physical form as a pre-requisite to qualify bidders.
5. Publishing bids on GeM for procurement of works.
6. Procurement of Goods by creating a Service bid on GeM & vice-versa.
7. Seeking sample with bid or approval of samples during bid evaluation process. However, trial / sample, as the case may be, shall be permitted in cases where trial / sample are allowed as per approved and published procurement policy of the Buyers' controlling Ministry / Department / State / Public Sector Enterprises Headquarters. If there is any violation of trial / sample clause with regard to approved policy of the Buyers' Ministry / Department / State / Public Sector Enterprises Headquarters, then this is to be determined and redressed by the concerned Buyer Organisation only.
8. Seeking experience from specific organization / department / institute only or from foreign / export experience.
9. Creating bid for items from incorrect categories.
10. Reference of conditions published on any external site or reference to external documents/clauses.
11. Asking for any Tender fee / Bid Participation fee, as the case may be.
12. Buyer added ATC Clauses which are in contravention of clauses defined in bid detail section, including specifications, EMD Detail, ePBG Detail and MII and MSE Purchase Preference sections of the bid, unless otherwise allowed by the applicable GeM GTC.
13. Any ATC clause in contravention with GeM GTC Clause 4 (xiii) (h) will be invalid.

In case of multiple L1 bidders against a service bid, the buyer shall place the Contract by selection of a bidder amongst the L-1 bidders through a Random Algorithm executed by GeM system.

14. In a category based bid, adding additional items, through buyer added, additional scope of work/ additional terms and conditions/or any other document. If buyer needs more items along with the main item, the same must be added through bunching category based items or by bunching custom catalogues or bunching a BoQ with the main category based item, the same must not be done through ATC or Scope of Work.

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

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

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

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

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

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

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

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

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

जेम की सामान्य शर्तों के खंड 26 के संदर्भ में भारत के साथ भूमि सीमा साझा करने वाले देश के बिडर से खरीद पर प्रतिबंध के संबंध में भारत के साथ भूमि सीमा साझा करने वाले देश का कोई भी बिडर इस निविदा में बिड देने के लिए तभी पात्र होगा जब वह बिड देने वाला सक्षम प्राधिकारी के पास पंजीकृत हो।बिड में भाग लेते समय बिडर को इसका अनुपालन करना होगा और कोई भी गलत घोषणा किए जाने व इसका अनुपालन न करने पर अनुबंध को तत्काल समाप्त करने और कानून के अनुसार आगे की कानूनी 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---