बिड संख्या/Bid Number: GEM/2025/B/6305388
दिनांक /Dated: 18-07-2025

बिड दस्तावेज़ / Bid Document

बिड विवरण/Bid Details	
बिड बंद होने की तारीख/समय /Bid End Date/Time	08-08-2025 17:00:00
बिड खुलने की तारीख/समय /Bid Opening Date/Time	08-08-2025 17:30:00
बिड पेशकश वैधता (बंद होने की तारीख से)/Bid Offer Validity (From End Date)	180 (Days)
मंत्रालय/राज्य का नाम/Ministry/State Name	Ministry Of Health And Family Welfare
विभाग का नाम/Department Name	Department Of Health And Family Welfare
संगठन का नाम/Organisation Name	All India Institute Of Medical Sciences (aiims)
कार्यालय का नाम/Office Name	Rishikesh
कुल मात्रा/Total Quantity	29993
वस्तु श्रेणी /Item Category	Blood Bags as per IS 15102 (Q2)
बिडर का न्यूनतम औसत वार्षिक टर्नओवर (3 वर्षों का) /Minimum Average Annual Turnover of the bidder (For 3 Years)	50 Lakh (s)
मूल उपकरण निर्माता का औसत टर्नओवर (गत 3 वर्षों का)/OEM Average Turnover (Last 3 Years)	100 Lakh (s)
उन्हीं/समान सेवा के लिए अपेक्षित विगत अनुभव के वर्ष/Years of Past Experience Required for same/similar service	1 Year (s)
वर्षों के अनुभव एवं टर्नओवर से एमएसई को छूट प्राप्त है / MSE Exemption for Years Of Experience and Turnover	Yes | Complete
स्टार्टअप के लिए अनुभव के वर्षों और टर्नओवर से छूट प्रदान की गई है / Startup Exemption for Years Of Experience and Turnover	Yes | Complete
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) *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 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 बढ़ाने के लिए आवश्यक न्यूनतम सहभागी विक्रेताओं की संख्या। / Minimum number of bids required to disable automatic bid extension	1
दिनों की संख्या, जिनके लिए बिड लगाने की समय-सीमा बढ़ाई जाएगी। / Number of days for which Bid would be auto-extended	7
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	Blood Bags as per IS 15102
तकनीकी मूल्यांकन के दौरान तकनीकी स्पष्टीकरण हेतु अनुमत समय /Time allowed for Technical Clarifications during technical evaluation	2 Days
निरीक्षण आवश्यक (सूचीबद्ध निरीक्षण प्राधिकरण /जेम के साथ पूर्व पंजीकृत एजेंसियों द्वारा)/Inspection Required (By Empanelled Inspection Authority / Agencies pre-registered with GeM)	No
अनुमानित बिड मूल्य /Estimated Bid Value	11377865
मूल्यांकन पद्धति/Evaluation Method	Item wise evaluation/
मध्यस्थता खंड/Arbitration Clause	No
सुलह खंड/Mediation Clause	No

ईएमडी विवरण/EMD Detail

एडवाईजरी बैंक/Advisory Bank	PUNJAB NATIONAL BANK
Schedule 1 ईएमडी राशि/EMD Amount (In INR)	101324
Schedule 2 ईएमडी राशि/EMD Amount (In INR)	236339
Schedule 3 ईएमडी राशि/EMD Amount (In INR)	3673

ईपीबीजी विवरण /ePBG Detail

एडवाइजरी बैंक/Advisory Bank	PUNJAB NATIONAL BANK
ईपीबीजी प्रतिशत (%)/ePBG Percentage(%)	5.00
ईपीबीजी की आवश्यक अवधि (माह) /Duration of ePBG required (Months).	15

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

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

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

लाभार्थी /Beneficiary :

RISHIKESH
Rishikesh, Department of Health and Family Welfare, All India Institute of Medical Sciences (AIIMS), Ministry of Health and Family Welfare
(Aiims)

विभाजन/Splitting

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

एमआईआई खरीद वरीयता/MII Purchase Preference

एमआईआई खरीद वरीयता/MII Purchase Preference	Yes

एमएसई खरीद वरीयता/MSE Purchase Preference

एमएसई खरीद वरीयता/MSE Purchase Preference	Yes

1. If the bidder is a Micro or Small Enterprise as per latest orders issued by Ministry of MSME, the bidder shall be exempted from the eligibility criteria of "Experience Criteria" as defined above subject to meeting of quality and technical specifications. The bidder seeking exemption from Experience Criteria, shall upload the supporting documents to prove his eligibility for exemption.
2. If the bidder is a Micro or Small Enterprise (MSE) as per latest orders issued by Ministry of MSME, the bidder shall be exempted 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 exempted from the "OEM Average Turnover" criteria also subject to meeting of quality and technical specifications. The bidder seeking exemption from Turnover, shall upload the supporting documents to prove his eligibility for exemption.
3. If the bidder is a DPIIT registered Startup, the bidder shall be exempted from the the eligibility criteria of "Experience Criteria" as defined above subject to their meeting of quality and technical specifications. The bidder seeking exemption from Experience Criteria, shall upload the supporting documents to prove his eligibility for

exemption.
4. If the bidder is a DPIIT registered Startup, the bidder shall be exempted 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 exempted from the "OEM Average Turnover" criteria also subject to meeting of quality and technical specifications. The bidder seeking exemption from Turnover shall upload the supporting documents to prove his eligibility for exemption.
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. 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.
11. 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.

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

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

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

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

मूल्यांकन अनुसूचियां / **Evaluation Schedules**	अनुमानित मूल्य / **Estimated Value**	वस्तु/श्रेणी / **Item/Category**	मात्रा / **Quantity**
Schedule 1	3377472	Blood Bags As Per Is 15102	8400
Schedule 2	7877953	Blood Bags As Per Is 15102	19593
Schedule 3	122440	Blood Bags As Per Is 15102	2000

Blood Bags As Per IS 15102 (8400 pieces)

(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)**
GENERAL FEATURES	Clinical Purpose	Collection, processing and storage of whole blood and blood components
	Disposable	Yes
PRODUCT INFORMATION	Conformity to standard for Blood Bag	ISO 3826/IS 15102: Latest Revision
	Type of blood bag	Quadruple (top and bottom)
	Capacity of blood Bag	350 ml
	Material of Bag (Medical grade)	DEHP Plasticized PVC

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
	Blood Collection Bags should be collapsible non vented sterile containers complete with collecting tube for completely closed system to avoid the chances of contamination	Yes
	Flexible pre-sterilized and pyrogen free	Yes
	Non toxic, non haemolytic, biocompatible material	Yes
	There should be no risk of contamination and air embolism (closed system) with all leak proof seals (Disposable bags)	Yes
	Slit on both sides of the bags should be enough to accommodate 5 -10 ml volume test tubes	Yes
	The capacity of the bag should be enough to prevent any bollooning/ripture of the abg from the seam when it is filled up with the requisite volume of blood	Yes
TUBING OF BAG	Flexible kink resistant tubing	Yes
	Non sticking	Yes
	Transparent	Yes
	Leak Proof	Yes
	Length of tubing from primary bag to needle	? 80 Cm
	The tubing should have same ID/segment number as that on the bag	Yes
	The tubes should have multiple printed ID/segment numbers	Yes
	Clamp provided for closed system	Yes
NEEDLE	Needle Size	16 G
	Ultra thin walled and straight to reduce penetration force and enable painless vein puncture	Yes
	Sharp, regular and smooth margins and bevelled tip	Yes
	Rust proof	Yes

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
	Tightly fixed with hub covered with sterile guard	Yes
	Hermetically sealed	Yes
	The needle should not separate from the tube at any point of time, especially while removing it from the vein for donor safety	Yes
	The needle must confirm to ISO 1135-3 standard	Yes
EXTERNAL PORT	Tamper proof and should not be re-capped	Yes
	Easily accessible	Yes
ANTICAOGULANT AND PRESERVATIVE SOLUTION	**Type of anticoagulant present**	CPD
	Quantity of anticoagulant solution	14 ml per 100 ml of blood
	Solution should be clear and colorless	Yes
	There should be no discoloration of solution on storage at room temperature	Yes
	Additive soultion present	Yes
	Type of additive solution	SAGM
	Quantity of Additive solution(ml)	80
	Anticoagulant and/or additive solution should be sterile and pyrogen free	Yes
	Availability of anticoagulant/additive quality check certificate from manufacture (proof of same to be submitted to buyer)	Yes
LABEL	Non-peel off	Yes
	Heat sealed/ Pressure embossed label	Yes
	The label should remain attached between room temperature to - 80°C with a tranparent adhesive	Yes
	Date of manufacturing, date of expiry and batch number must be mentioned on each bag	Yes

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
RESISTANCE TO DISTORTION	Bag (Fiiled to normal capacity) shall withstand a acceleration of 5000 g for 30 min at temperature 4°C to 24°C without becoming permanently distorted	Yes
	Bag (Fiiled to normal capacity) should be able to withstand temperature upto - 80°C without breakage	Yes
PACKAGING	Individual bag packed in plastic pack and multiple bags packed in moisture proof aluminum foil (Protective dual packaging) eliminating microbial contamination on surface maintaining the contents of the bag	Yes
	The supplier should ensure proper transportation of the consignment of blood bags in temperature controlled conditions (Storage temperature should not exceed 30°C)	Yes
CERTIFICATIONS & REPORTS	**Manufacturer certifications (Proof of the same to be submitted to the buyer on demand)**	ISO 13485, GMP, WHO GMP
	Product Certifications (Proof of the same to be submitted to buyer on demand)	ISO, EU-CE
	Each batch supplied should be accompanied with quality assurance test report from NABL approved lab/any lab approved fron govt of India as well as in house lab	Yes
	Biocompatibility of the material of the product must certified by the manufacturer and be supported by the test reports of cell culture cytotoxicity, hemolysis, systemic infections, sensitization, Inra-cutaneous injection, pyrogen test and Sterility	Yes

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
	Submission of manufacturer's documented evidence of biochemical parameters of blood stored in CPDA/CPDA-1/CPD-SAGM containing DEHP plasticized PVC blood bags manufactured by the company on 28th/35th/42nd day of storage	Yes
SHELF LIFE	**Shelf Life from the date of manufacture (in months)**	36, 24
	Stability report from a recogniozed laboratory must be submitted to the buyer at the time of supply	Yes
	The product should have atleast 3/4 of the total shelf life at the time of dispatch to the consignee	Yes

Additional Specification Parameters - Blood Bags As Per IS 15102 (8400 pieces)

Specification Parameter Name	Bid Requirement (Allowed Values)
Notwithstanding anything stated above, the technical specifications mentioned in the ATC must be complied with.	Notwithstanding anything stated above, the technical specifications mentioned in the ATC must be complied with.

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

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

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

क्र.सं./S.No.	परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer	पता/Address	डिलीवरी अनुसूची /Delivery Schedule अनुबंध prarambh होने की तारीख से दिनों की संख्या में /(In number of days from contract start days)
1	RAJEEV KUMAR	249203,Central Store, AIIMS Rishikesh, Virbhadra Road, Near Bairaj Lake, Rishikesh, Uttarakhand-249203	

मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____तक पूरी कर ली जाए /Delivery to be completed by
2100	1	30
2100	91	105
2100	181	195
2100	271	285

Blood Bags As Per IS 15102 (19593 pieces)

(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)
GENERAL FEATURES	Clinical Purpose	Collection, processing and storage of whole blood and blood components
	Disposable	Yes
PRODUCT INFORMATION	Conformity to standard for Blood Bag	ISO 3826/IS 15102: Latest Revision
	Type of blood bag	Quadruple (top and bottom)
	Capacity of blood Bag	450 ml
	Material of Bag (Medical grade)	DEHP Plasticized PVC
	Blood Collection Bags should be collapsible non vented sterile containers complete with collecting tube for completely closed system to avoid the chances of contamination	Yes
	Flexible pre-sterilized and pyrogen free	Yes
	Non toxic, non haemolytic, biocompatible material	Yes

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
	There should be no risk of contamination and air embolism (closed system) with all leak proof seals (Disposable bags)	Yes
	Slit on both sides of the bags should be enough to accommodate 5 -10 ml volume test tubes	Yes
	The capacity of the bag should be enough to prevent any bollooning/ripture of the abg from the seam when it is filled up with the requisite volume of blood	Yes
TUBING OF BAG	Flexible kink resistant tubing	Yes
	Non sticking	Yes
	Transparent	Yes
	Leak Proof	Yes
	Length of tubing from primary bag to needle	? 80 Cm
	The tubing should have same ID/segment number as that on the bag	Yes
	The tubes should have multiple printed ID/segment numbers	Yes
	Clamp provided for closed system	Yes
NEEDLE	Needle Size	16 G
	Ultra thin walled and straight to reduce penetration force and enable painless vein puncture	Yes
	Sharp, regular and smooth margins and bevelled tip	Yes
	Rust proof	Yes
	Tightly fixed with hub covered with sterile guard	Yes
	Hermetically sealed	Yes
	The needle should not separate from the tube at any point of time, especially while removing it from the vein for donor safety	Yes
	The needle must confirm to ISO 1135-3 standard	Yes

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
EXTERNAL PORT	Tamper proof and should not be re-capped	Yes
	Easily accessible	Yes
ANTICAOGULANT AND PRESERVATIVE SOLUTION	**Type of anticoagulant present**	CPD
	Quantity of anticoagulant solution	14 ml per 100 ml of blood
	Solution should be clear and colorless	Yes
	There should be no discoloration of solution on storage at room temperature	Yes
	Additive soultion present	Yes
	Type of additive solution	SAGM
	Quantity of Additive solution(ml)	100
	Anticoagulant and/or additive solution should be sterile and pyrogen free	Yes
	Availability of anticoagulant/additive quality check certificate from manufacture (proof of same to be submitted to buyer)	Yes
LABEL	Non-peel off	Yes
	Heat sealed/ Pressure embossed label	Yes
	The label should remain attached between room temperature to - 80°C with a tranparent adhesive	Yes
	Date of manufacturing, date of expiry and batch number must be mentioned on each bag	Yes
RESISTANCE TO DISTORTION	Bag (Fiiled to normal capacity) shall withstand a acceleration of 5000 g for 30 min at temperature 4°C to 24°C without becoming permanently distorted	Yes
	Bag (Fiiled to normal capacity) should be able to withstand temperature upto - 80°C without breakage	Yes

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
PACKAGING	Individual bag packed in plastic pack and multiple bags packed in moisture proof aluminum foil (Protective dual packaging) eliminating microbial contamination on surface maintaining the contents of the bag	Yes
	The supplier should ensure proper transportation of the consignment of blood bags in temperature controlled conditions (Storage temperature should not exceed 30°C)	Yes
CERTIFICATIONS & REPORTS	**Manufacturer certifications (Proof of the same to be submitted to the buyer on demand)**	ISO 13485, GMP, WHO GMP
	Product Certifications (Proof of the same to be submitted to buyer on demand)	ISO, BIS, EU-CE, US-FDA
	Each batch supplied should be accompanied with quality assurance test report from NABL approved lab/any lab approved fron govt of India as well as in house lab	Yes
	Biocompatibility of the material of the product must certified by the manufacturer and be supported by the test reports of cell culture cytotoxicity, hemolysis, systemic infections, sensitization, Inra-cutaneous injection, pyrogen test and Sterility	Yes
	Submission of manufacturer's documented evidence of biochemical parameters of blood stored in CPDA/CPDA-1/CPD-SAGM containing DEHP plasticized PVC blood bags manufactured by the company on 28th/35th/42nd day of storage	Yes
SHELF LIFE	**Shelf Life from the date of manufacture (in months)**	36, 24
	Stability report from a recogniozed laboratory must be submitted to the buyer at the time of supply	Yes

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
	The product should have atleast 3/4 of the total shelf life at the time of dispatch to the consignee	Yes

Additional Specification Parameters - Blood Bags As Per IS 15102 (19593 pieces)

Specification Parameter Name	Bid Requirement (Allowed Values)
Notwithstanding anything stated above, the technical specifications mentioned in the ATC must be complied with.	Notwithstanding anything stated above, the technical specifications mentioned in the ATC must be complied with.

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

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

क्र.सं./S.No.	परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer	पता/Address	डिलीवरी अनुसूची /Delivery Schedule अनुबंध prarambh होने की तारीख से दिनों की संख्या में /(In number of days from contract start days)		
			मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____तक पूरी कर ली जाए /Delivery to be completed by
1	RAJEEV KUMAR	249203,Central Store, AIIMS Rishikesh, Virbhadra Road, Near Bairaj Lake, Rishikesh, Uttarakhand-249203	4900	1	30
			4900	91	105
			4900	181	195
			4893	271	285

Blood Bags As Per IS 15102 (2000 pieces)

(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)
GENERAL FEATURES	Clinical Purpose	Collection, processing and storage of whole blood and blood components
	Disposable	Yes
PRODUCT INFORMATION	Conformity to standard for Blood Bag	ISO 3826/IS 15102: Latest Revision
	Type of blood bag	Transfer bag system
	Capacity of blood Bag	300 ml
	Material of Bag (Medical grade)	DEHP Plasticized PVC
	Blood Collection Bags should be collapsible non vented sterile containers complete with collecting tube for completely closed system to avoid the chances of contamination	Yes
	Flexible pre-sterilized and pyrogen free	Yes
	Non toxic, non haemolytic, biocompatible material	Yes
	There should be no risk of contamination and air embolism (closed system) with all leak proof seals (Disposable bags)	Yes
	Slit on both sides of the bags should be enough to accommodate 5 -10 ml volume test tubes	Yes
	The capacity of the bag should be enough to prevent any bollooning/ripture of the abg from the seam when it is filled up with the requisite volume of blood	Yes
TUBING OF BAG	Flexible kink resistant tubing	Yes
	Non sticking	Yes
	Transparent	Yes
	Leak Proof	Yes
	Length of tubing from primary bag to needle	? 80 Cm
	The tubing should have same ID/segment number as that on the bag	Yes
	The tubes should have multiple printed ID/segment numbers	Yes

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
	Clamp provided for closed system	Yes
NEEDLE	Needle Size	16 G
	Ultra thin walled and straight to reduce penetration force and enable painless vein puncture	Yes
	Sharp, regular and smooth margins and bevelled tip	Yes
	Rust proof	Yes
	Tightly fixed with hub covered with sterile guard	Yes
	Hermetically sealed	Yes
	The needle should not separate from the tube at any point of time, especially while removing it from the vein for donor safety	Yes
	The needle must confirm to ISO 1135-3 standard	Yes
EXTERNAL PORT	Tamper proof and should not be re-capped	Yes
	Easily accessible	Yes
ANTICAOGULANT AND PRESERVATIVE SOLUTION	**Type of anticoagulant present**	CPDA-1
	Quantity of anticoagulant solution	14 ml per 100 ml of blood
	Solution should be clear and colorless	Yes
	There should be no discoloration of solution on storage at room temperature	Yes
	Additive soultion present	No
	Type of additive solution	NA
	Quantity of Additive solution(ml)	NA
	Anticoagulant and/or additive solution should be sterile and pyrogen free	Yes
	Availability of anticoagulant/additive quality check certificate from manufacture (proof of same to be submitted to buyer)	Yes
LABEL	Non-peel off	Yes

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
	Heat sealed/ Pressure embossed label	Yes
	The label should remain attached between room temperature to - 80°C with a tranparent adhesive	Yes
	Date of manufacturing, date of expiry and batch number must be mentioned on each bag	Yes
RESISTANCE TO DISTORTION	Bag (Fiiled to normal capacity) shall withstand a acceleration of 5000 g for 30 min at temperature 4°C to 24°C without becoming permanently distorted	Yes
	Bag (Fiiled to normal capacity) should be able to withstand temperature upto - 80°C without breakage	Yes
PACKAGING	Individual bag packed in plastic pack and multiple bags packed in moisture proof aluminum foil (Protective dual packaging) eliminating microbial contamination on surface maintaining the contents of the bag	Yes
	The supplier should ensure proper transportation of the consignment of blood bags in temperature controlled conditions (Storage temperature should not exceed 30°C)	Yes
CERTIFICATIONS & REPORTS	**Manufacturer certifications (Proof of the same to be submitted to the buyer on demand)**	ISO 13485, GMP, WHO GMP
	Product Certifications (Proof of the same to be submitted to buyer on demand)	ISO, EU-CE, US-FDA
	Each batch supplied should be accompanied with quality assurance test report from NABL approved lab/any lab approved fron govt of India as well as in house lab	Yes

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
	Biocompatibility of the material of the product must certified by the manufacturer and be supported by the test reports of cell culture cytotoxicity, hemolysis, systemic infections, sensitization, Inra-cutaneous injection, pyrogen test and Sterility	Yes
	Submission of manufacturer's documented evidence of biochemical parameters of blood stored in CPDA/CPDA-1/CPD-SAGM containing DEHP plasticized PVC blood bags manufactured by the company on 28th/35th/42nd day of storage	Yes
SHELF LIFE	**Shelf Life from the date of manufacture (in months)**	24
	Stability report from a recogniozed laboratory must be submitted to the buyer at the time of supply	Yes
	The product should have atleast 3/4 of the total shelf life at the time of dispatch to the consignee	Yes

Additional Specification Parameters - Blood Bags As Per IS 15102 (2000 pieces)

Specification Parameter Name	Bid Requirement (Allowed Values)
Notwithstanding anything stated above, the technical specifications mentioned in the ATC must be complied with.	Notwithstanding anything stated above, the technical specifications mentioned in the ATC must be complied with.

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

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

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

<table>
<tr><th>क्र.सं./S.No.</th><th>परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer</th><th>पता/Address</th><th>डिलीवरी अनुसूची /Delivery Schedule अनुबंध prarambh होने की तारीख से दिनों की संख्या में /(In number of days from contract start days)</th></tr>
<tr><td>1</td><td>RAJEEV KUMAR</td><td>249203,Central Store, AIIMS Rishikesh, Virbhadra Road, Near Bairaj Lake, Rishikesh, Uttarakhand-249203</td><td>
<table>
<tr><th>मात्रा /Quantity</th><th>प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after</th><th>डिलीवरी _____तक पूरी कर ली जाए /Delivery to be completed by</th></tr>
<tr><td>500</td><td>1</td><td>30</td></tr>
<tr><td>500</td><td>91</td><td>105</td></tr>
<tr><td>500</td><td>181</td><td>195</td></tr>
<tr><td>500</td><td>271</td><td>285</td></tr>
</table>
</td></tr>
</table>

Special terms and conditions-Version:2 effective from 20-10-2022 for category Blood Bags as per IS 15102

1. 1. All Provisions of Drugs and Cosmetics Act, 1940 and Rules made there under as amended till date will always be applicable. This will include all notifications issued by *Central Drugs Standard Control Organisation* (*CDSCO*), Ministry of Health & Family Welfare (MoHFW) and Department of Pharmaceuticals (DOP), Ministry of Chemicals & Fertilizers time to time in this regard.
   2. The sellers are registered on GeM based on self-declaration of valid Drug License, product certification, test reports etc. However, buyers must check and validate the details at their end for all applicable licenses and certifications e.g., validity and authenticity/genuineness of drug license, product certification, manufacturer certification/licenses, test reports etc.
   3. The price offered by the seller shall not, in any case exceed the DPCO controlled price, if any, fixed by the Central/State Government, the Maximum Retail Price (MRP) and the selling price. The seller must reduce the prices if there is any reduction in DPCO ceiling price, if any.
   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 in the bid to ensure items are procured from authentic/validated source with appropriate and applicable quality. The above terms and conditions are in reverse order of precedence i.e., ATC shall supersede specific STC which shall supersede General Terms and Conditions (“GTC”), whenever there are any conflicting provisions.

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

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

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

5. **Turnover**

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

6. **Sample Clause**

   For clothing and allied items, if pre-dispatch Inspection clause has been selected in the Bid, the Inspection Agency shall forward sample from the accepted lot duly identified/ sealed by it, as Reference Sample to each consignee (one reference sample per consignee) for comparing the lot received at consignee end with such reference sample. Such reference samples will be treated as part of supplied quantity from the lot and cost shall be borne by the Buyer.

7. **Forms of EMD and PBG**

   Successful Bidder can submit the Performance Security in the form of Fixed Deposit Receipt also (besides PBG which is allowed as per GeM GTC). FDR should be made out or pledged in the name of

   AIIMS RISHIKESH
   A/C (Name of the Seller). The bank should certify on it that the deposit can be withdrawn only on the demand or with the sanction of the pledgee. For release of Security Deposit, the FDR will be released in favour of bidder by the Buyer after making endorsement on the back of the FDR duly signed and stamped along with covering letter. Successful Bidder has to upload scanned copy of the FDR document in place of PBG and has to ensure delivery of hard copy of Original FDR to the Buyer within 15 days of award of contract.

8. **Buyer Added Bid Specific ATC**

   Buyer uploaded ATC document Click here to view the file.

अस्वीकरण/Disclaimer

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

1. Definition of Class I and Class II suppliers in the bid not in line with the extant Order / Office Memorandum issued by DPIIT in this regard.
2. Seeking EMD submission from bidder(s), including via Additional Terms & Conditions, in contravention to exemption provided to such sellers under GeM GTC.
3. Publishing Custom / BOQ bids for items for which regular GeM categories are available without any Category item bunched with it.
4. Creating BoQ bid for single item.
5. Mentioning specific Brand or Make or Model or Manufacturer or Dealer name.
6. Mandating submission of documents in physical form as a pre-requisite to qualify bidders.
7. Floating / creation of work contracts as Custom Bids in Services.
8. Seeking sample with bid or approval of samples during bid evaluation process. (However, in bids for attached categories, trials are allowed as per approved procurement policy of the buyer nodal Ministries)
9. Mandating foreign / international certifications even in case of existence of Indian Standards without specifying equivalent Indian Certification / standards.

10. Seeking experience from specific organization / department / institute only or from foreign / export experience.
11. Creating bid for items from irrelevant categories.
12. Incorporating any clause against the MSME policy and Preference to Make in India Policy.
13. Reference of conditions published on any external site or reference to external documents/clauses.
14. Asking for any Tender fee / Bid Participation fee / Auction fee in case of Bids / Forward Auction, as the case may be.
15. Buyer added ATC Clauses which are in contravention of clauses defined by buyer in system generated bid template as indicated above in the Bid Details section, EMD Detail, ePBG Detail and MII and MSE Purchase Preference sections of the bid, unless otherwise allowed by GeM GTC.
16. In a category based bid, adding additional items, through buyer added additional scope of work/ additional terms and conditions/or any other document. If buyer needs more items along with the main item, the same must be added through bunching category based items or by bunching custom catalogs or bunching a BoQ with the main category based item, the same must not be done through ATC or Scope of Work.

Further, if any seller has any objection/grievance against these additional clauses or otherwise on any aspect of this bid, they can raise their representation against the same by using the Representation window provided in the bid details field in Seller dashboard after logging in as a seller within 4 days of bid publication on GeM. Buyer is duty bound to reply to all such representations and would not be allowed to open bids if he fails to reply to such representations.

All GeM Sellers / Service Providers are mandated to ensure compliance with all the applicable laws / acts / rules including but not limited to all Labour Laws 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. Any non-compliance will be treated as breach of contract and Buyer may take suitable actions as per GeM Contract.

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---