

बिड दस्तावेज़ / Bid Document

बिड विवरण / Bid Details	
बिड बंद होने की तारीख/समय / Bid End Date/Time	09-05-2026 17:00:00
बिड खुलने की तारीख/समय / Bid Opening Date/Time	09-05-2026 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 Labour And Employment
विभाग का नाम / Department Name	Na
संगठन का नाम / Organisation Name	Employees State Insurance Corporation (esic)
कार्यालय का नाम / Office Name	Jammu And Kashmir
कुल मात्रा / Total Quantity	72885
वस्तु श्रेणी / Item Category	Rapid Test Kit for HIV / Syphilis / Hepatitis B Virus (HBV) / Hepatitis C Virus (HCV) (Q2) , Point of Care Rapid Test Kits for Humans - Dengue, Malaria, Typhoid & Others (Q2) , Rapid Pregnancy Test Kit (Q2) , Blood Collection Tube (Q2) , Blood Collection Needles (V2) (Q2)
बिडर का न्यूनतम औसत वार्षिक टर्नओवर (3 वर्षों का) / Minimum Average Annual Turnover of the bidder (For 3 Years)	4 Lakh (s)
मूल उपकरण निर्माता का औसत टर्नओवर (गत 3 वर्षों का) / OEM Average Turnover (Last 3 Years)	18 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 Complete
स्टार्टअप के लिए अनुभव के वर्षों और टर्नओवर से छूट प्रदान की गई है / Startup Relaxation for Years Of Experience and Turnover	Yes Complete
विक्रेता से मांगे गए दस्तावेज़ / Document required from seller	Experience Criteria, Past Performance, Bidder Turnover, 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 दस्तावेजों 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	3
विगत प्रदर्शन /Past Performance	50 %
बिड से रिवर्स नीलामी सक्रिय किया/Bid to RA enabled	No
बिड का प्रकार/Type of Bid	Two Packet Bid
प्राथमिक उत्पाद श्रेणी/Primary product category	Point of Care Rapid Test Kits for Humans - Dengue, Malaria, Typhoid & Others
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	16000

ईपीबीजी विवरण /ePBG Detail

एडवाइजरी बैंक/Advisory Bank	State Bank of India
ईपीबीजी प्रतिशत (%) /ePBG Percentage(%)	3.00

ईपीबीजी की आवश्यक अवधि (माह) /Duration of ePBG required (Months).

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(a). जेम की शर्तों के अनुसार ईएमडी छूट के इच्छुक बिडर को संबंधित कटेगरी के लिए बिड के साथ वैध समर्थित दस्तावेज 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 Model Hospital, Lane no 2 , SIDCO industrial Area, Bari Brahmana, Jammu, Jammu And Kashmir 181133
(Medical Superintendent)

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

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.

Pre Bid Detail(s)

मूल्य भिन्नता खंड दस्तावेज़/Pre-Bid Date and Time	प्री-बिड स्थान/Pre-Bid Venue
30-04-2026 11:00:00	Conferance Hall M S Office, ESIC Moel hospital, bari brahmana, jammu

Rapid Test Kit For HIV / Syphilis / Hepatitis B Virus (HBV) / Hepatitis C Virus (HCV) (100 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	Type of Test Kit	HIV1 & HIV2 Antibodies Detection Rapid Test Kit
	Detection Type	Qualitative
	Testing Method	Lateral Flow (Immunochromatography)
	Result Time	? 15 minutes
	Positive and negative controls provided with each pack of kit	Yes
PACKAGING	Number of Tests per Pack	50 Tests
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

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 प्रारम्भ होने की तारीख से दिनों की संख्या में / (In number of days from contract start days)		
1	Rakesh Choudhary	181133,ESIC MODEL HOSPITAL, LANE NO.-02 - SIDCO, OPPOSITE POLICE STATION, BARI BRAHMANA JAMMU	मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____तक पूरी कर ली जाए /Delivery to be completed by
			35	1	15
			35	105	135
			30	210	240

Rapid Test Kit For HIV / Syphilis / Hepatitis B Virus (HBV) / Hepatitis C Virus (HCV) (100 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	Type of Test Kit	Hepatitis C Virus (HCV) Total Antibodies Detection Rapid Test Kit
	Detection Type	Qualitative
	Result Time	? 15 minutes
	Positive and negative controls provided with each pack of kit	Yes
PACKAGING	Number of Tests per Pack	50 Tests

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	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

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

क्र.सं./S.N o.	परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer	पता/Address	डिलीवरी अनुसूची /Delivery Schedule अनुबंध प्रारम्भ होने की तारीख से दिनों की संख्या में / (In number of days from contract start days)		
1	Rakesh Choudhary	181133,ESIC MODEL HOSPITAL, LANE NO.-02 - SIDCO, OPPOSITE POLICE STATION, BARI BRAHMANA JAMMU	मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____ तक पूरी कर ली जाए /Delivery to be completed by
			35	1	15
			35	105	135
			30	210	255

Rapid Test Kit For HIV / Syphilis / Hepatitis B Virus (HBV) / Hepatitis C Virus (HCV) (100 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	Type of Test Kit	Hepatitis B Surface Antigen (HBsAg) Rapid Test Kit
	Detection Type	Qualitative

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
	Testing Method	Lateral Flow (Immunochromatography)
	Result Time	? 15 minutes
	Positive and negative controls provided with each pack of kit	Yes
PACKAGING	Number of Tests per Pack	50 Tests
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

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

क्र.सं./S.N o.	परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer	पता/Address	डिलीवरी अनुसूची /Delivery Schedule अनुबंध प्रारंभ होने की तारीख से दिनों की संख्या में / (In number of days from contract start days)		
1	Rakesh Choudhary	181133,ESIC MODEL HOSPITAL, LANE NO.-02 - SIDCO, OPPOSITE POLICE STATION, BARI BRAHMANA JAMMU	मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____तक पूरी कर ली जाए /Delivery to be completed by
			35	1	15
			35	105	135
			30	210	255

Rapid Test Kit For HIV / Syphilis / Hepatitis B Virus (HBV) / Hepatitis C Virus (HCV) (20 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	Type of Test Kit	HIV1 & HIV2 Antibodies Detection Rapid Test Kit, HIV1 & HIV2 Antibodies and HIV1 p24 Antigen Detection Rapid Test Kit, HIV1 (all subtypes) , HIV2 and Syphilis (Treponema pallidum) Antibody Combo Rapid Test Kit, Hepatitis C Virus (HCV) Total Antibodies Detection Rapid Test Kit, Hepatitis C Virus (HCV) Antibody and Antigen Detection Rapid Test Kit, Hepatitis B Surface Antigen (HBsAg) Rapid Test Kit, HIV 1/2 Antibodies + Hepatitis C Virus (HCV) Antibodies + Syphilis Antibodies + Hepatitis B Surface Antigen (HBsAg) Combo Rapid Test Kit
	Detection Type	Qualitative
	Testing Method	Flow-Through (Immunoconcentration), Lateral Flow (Immunochromatography), Agglutination, Dot Immuno Assay
	Result Time	? 15 minutes
	Positive and negative controls provided with each pack of kit	Yes
PACKAGING	Number of Tests per Pack	50 Tests
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

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

क्र.सं./S.N o.	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.	प्रेषित/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer	पता/Address	डिलीवरी अनुसूची /Delivery Schedule अनुबंध प्रारम्भ होने की तारीख से दिनों की संख्या में / (In number of days from contract start days)		
1	Rakesh Choudhary	181133,ESIC MODEL HOSPITAL, LANE NO.-02 - SIDCO, OPPOSITE POLICE STATION, BARI BRAHMANA JAMMU	मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____ तक पूरी कर ली जाए /Delivery to be completed by
			7	1	15
			7	105	135
			6	210	255

Point Of Care Rapid Test Kits For Humans - Dengue, Malaria, Typhoid & Others (30 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	Type of Test Kit	Malaria Plasmodium falciparum (Pf) and Plasmodium vivax (Pv) Antigen Rapid Test Kit
	Detection Type	Qualitative
	Sample Type	Whole Blood, Serum and Plasma
	Result Time	? 15 minutes
	Positive and negative controls provided with each pack of kit	Yes
PACKAGING	Number of Tests per Pack	50 Tests
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

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

क्र.सं./S.N o.	परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer	पता/Address	डिलीवरी अनुसूची /Delivery Schedule अनुबंध प्रारंभ होने की तारीख से दिनों की संख्या में /(In number of days from contract start days)		
1	Rakesh Choudhary	181133,ESIC MODEL HOSPITAL, LANE NO.-02 - SIDCO, OPPOSITE POLICE STATION, BARI BRAHMANA JAMMU	मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____तक पूरी कर ली जाए /Delivery to be completed by
			10	1	15
			10	105	135
			10	210	255

Point Of Care Rapid Test Kits For Humans - Dengue, Malaria, Typhoid & Others (200 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	Type of Test Kit	Chikungunya IgM and IgG Antibodies Detection Rapid Test Kit
	Detection Type	Qualitative
	Sample Type	Whole Blood, Serum and Plasma
	Result Time	? 15 minutes
	Positive and negative controls provided with each pack of kit	Yes
PACKAGING	Number of Tests per Pack	10 Tests
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

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 प्रारम्भ होने की तारीख से दिनों की संख्या में / (In number of days from contract start days)		
1	Rakesh Choudhary	181133,ESIC MODEL HOSPITAL, LANE NO.-02 - SIDCO, OPPOSITE POLICE STATION, BARI BRAHMANA JAMMU	मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____तक पूरी कर ली जाए /Delivery to be completed by
			70	1	15
			70	105	135
			60	210	240

Point Of Care Rapid Test Kits For Humans - Dengue, Malaria, Typhoid & Others (150 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	Type of Test Kit	Typhoid - Salmonella Typhi IgM and IgG Antibodies Detection Rapid Test Kit
	Detection Type	Qualitative
	Sample Type	Whole Blood, Serum and Plasma
	Result Time	? 15 minutes
	Sensitivity (%)	? 99 Or higher
	Specificity (%)	? 99 Or higher

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
	Positive and negative controls provided with each pack of kit	Yes
PACKAGING	Number of Tests per Pack	10 Tests
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

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 प्रारम्भ होने की तारीख से दिनों की संख्या में /((In number of days from contract start days)		
1	Rakesh Choudhary	181133,ESIC MODEL HOSPITAL, LANE NO.-02 - SIDCO, OPPOSITE POLICE STATION, BARI BRAHMANA JAMMU	मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____तक पूरी कर ली जाए /Delivery to be completed by
			50	1	15
			50	105	135
			50	210	240

Point Of Care Rapid Test Kits For Humans - Dengue, Malaria, Typhoid & Others (100 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	Type of Test Kit	Dengue NS1 Antigen and IgM + IgG Antibodies Detection Rapid Test Kit
	Detection Type	Qualitative
	Sample Type	Whole Blood, Serum and Plasma
	Result Time	? 15 minutes
	Sensitivity (%)	? 98, ? 99 Or higher
	Specificity (%)	? 98, ? 99 Or higher
	Positive and negative controls provided with each pack of kit	Yes
PACKAGING	Number of Tests per Pack	10 Tests
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

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 प्रारंभ 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	Rakesh Choudhary	181133,ESIC MODEL HOSPITAL, LANE NO.-02 - SIDCO, OPPOSITE POLICE STATION, BARI BRAHMANA JAMMU			
			35	1	15
			35	105	135
			30	210	245

Point Of Care Rapid Test Kits For Humans - Dengue, Malaria, Typhoid & Others (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	Type of Test Kit	Fecal Occult Blood (FOB) Rapid Test Kit
	Detection Type	Qualitative
	Sample Type	Stool (Feces)
	Result Time	? 15 minutes
	Sensitivity (%)	? 99 Or higher
	Specificity (%)	? 99 Or higher
	Positive and negative controls provided with each pack of kit	Yes, No
PACKAGING	Number of Tests per Pack	10 Tests
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

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

क्र.सं./S.N o.	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)
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क्र.सं./S.No.	परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer	पता/Address	डिलीवरी अनुसूची /Delivery Schedule अनुबंध प्रारम्भ होने की तारीख से दिनों की संख्या में /In number of days from contract start days)									
1	Rakesh Choudhary	181133,ESIC MODEL HOSPITAL, LANE NO.-02 - SIDCO, OPPOSITE POLICE STATION, BARI BRAHMANA JAMMU	<table border="1"> <thead> <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> </thead> <tbody> <tr> <td>3</td> <td>1</td> <td>15</td> </tr> <tr> <td>2</td> <td>150</td> <td>165</td> </tr> </tbody> </table>	मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____तक पूरी कर ली जाए /Delivery to be completed by	3	1	15	2	150	165
मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____तक पूरी कर ली जाए /Delivery to be completed by										
3	1	15										
2	150	165										

Rapid Pregnancy Test Kit (80 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	Detection Type	Qualitative
	Sample Type	Urine
	Sensitivity	?25 milli I.U/ ml
	Specificity	?99%
PACKAGING	Number of Tests per Pack	25 Tests
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

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 प्रारंभ होने की तारीख से दिनों की संख्या में / (In number of days from contract start days)		
1	Rakesh Choudhary	181133,ESIC MODEL HOSPITAL, LANE NO.-02 - SIDCO, OPPOSITE POLICE STATION, BARI BRAHMANA JAMMU	मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____तक पूरी कर ली जाए /Delivery to be completed by
			30	1	15
			25	105	135
			25	210	240

Blood Collection Tube (5000 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	Product Description	Blood Collection Tube
	Usage	Single-Use (Disposable)
	Sterility	Sterile
TECHNICAL INFORMATION	Type of Tube	Vacuum Blood Collection Tube (Evacuated Tube)
	Material of Tube	Polyethylene Terephthalate (PET)
	Non toxic, leakproof and crack resistance material	Yes
	Transparency	Transparent
	Negative air pressure present inside the tube	Yes
	Additive	3.8% Sodium Citrate
	Draw volume of blood sample	2 mL
	Cap Color	Black
	Patient Utility	Adult

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
	Graduation marking	Yes
	Label on tube for Identification	Yes
CERTIFICATIONS	Compliance to Medical Device Rules (MDR) 2017 as amended till date	Yes
	Availability of valid medical device license for the product issued from the competent authority defined under Drugs and Cosmetic Act 1940 and Rules made there under as amended till date	Yes
	Manufacturer Certification	ISO:13485 (Latest)
	Availability of Test Report for each supplied batch/product as per Medical Device Rules (MDR) 2017 as amended till date	Yes
	Submission of all necessary certifications, licenses and test reports to the buyer at the time of bid submission and/or along with supplies as per buyer requirement	Yes
SHELF LIFE	Minimum shelf life of the product at the time of delivery to the consignee	3/4 th of Total Shelf Life
ADVANCE SAMPLE	Agree to provide advance sample of the product for buyer's approval after award of contract and before commencement of supply in case of bidding only	Yes

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

क्र.सं./S.N o.	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 प्रारम्भ होने की तारीख से दिनों की संख्या में /((In number of days from contract start days)												
1	Rakesh Choudhary	181133,ESIC MODEL HOSPITAL, LANE NO.-02 - SIDCO, OPPOSITE POLICE STATION, BARI BRAHMANA JAMMU	<table border="1"> <thead> <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> </thead> <tbody> <tr> <td>1800</td> <td>1</td> <td>15</td> </tr> <tr> <td>1800</td> <td>105</td> <td>135</td> </tr> <tr> <td>1400</td> <td>210</td> <td>240</td> </tr> </tbody> </table>	मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____तक पूरी कर ली जाए /Delivery to be completed by	1800	1	15	1800	105	135	1400	210	240
मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____तक पूरी कर ली जाए /Delivery to be completed by													
1800	1	15													
1800	105	135													
1400	210	240													

Blood Collection Tube (5000 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 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)
GENERAL	Product Description	Blood Collection Tube
	Usage	Single-Use (Disposable)
	Sterility	Sterile
TECHNICAL INFORMATION	Type of Tube	Vacuum Blood Collection Tube (Evacuated Tube)
	Material of Tube	Polyethylene Terephthalate (PET)
	Non toxic, leakproof and crack resistance material	Yes
	Transparency	Transparent
	Negative air pressure present inside the tube	Yes
	Additive	3.2% Sodium Citrate
	Draw volume of blood sample	1.8 mL
Cap Color	Light Blue	

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
	Patient Utility	Adult
	Graduation marking	Yes
	Label on tube for Identification	Yes
CERTIFICATIONS	Compliance to Medical Device Rules (MDR) 2017 as amended till date	Yes
	Availability of valid medical device license for the product issued from the competent authority defined under Drugs and Cosmetic Act 1940 and Rules made there under as amended till date	Yes
	Manufacturer Certification	ISO:13485 (Latest)
	Availability of Test Report for each supplied batch/product as per Medical Device Rules (MDR) 2017 as amended till date	Yes
	Submission of all necessary certifications, licenses and test reports to the buyer at the time of bid submission and/or along with supplies as per buyer requirement	Yes
SHELF LIFE	Minimum shelf life of the product at the time of delivery to the consignee	3/4 th of Total Shelf Life
ADVANCE SAMPLE	Agree to provide advance sample of the product for buyer's approval after award of contract and before commencement of supply in case of bidding only	Yes

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

क्र.सं./S.N o.	परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer	पता/Address	डिलीवरी अनुसूची /Delivery Schedule अनुबंध प्रारम्भ होने की तारीख से दिनों की संख्या में /(In number of days from contract start days)

क्र.सं./S.No.	प्रेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer	पता/Address	डिलीवरी अनुसूची /Delivery Schedule अनुबंध प्रारम्भ होने की तारीख से दिनों की संख्या में / (In number of days from contract start days)												
1	Rakesh Choudhary	181133,ESIC MODEL HOSPITAL, LANE NO.-02 - SIDCO, OPPOSITE POLICE STATION, BARI BRAHMANA JAMMU	<table border="1"> <thead> <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> </thead> <tbody> <tr> <td>1800</td> <td>1</td> <td>15</td> </tr> <tr> <td>1800</td> <td>105</td> <td>135</td> </tr> <tr> <td>1400</td> <td>210</td> <td>245</td> </tr> </tbody> </table>	मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____ तक पूरी कर ली जाए /Delivery to be completed by	1800	1	15	1800	105	135	1400	210	245
मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____ तक पूरी कर ली जाए /Delivery to be completed by													
1800	1	15													
1800	105	135													
1400	210	245													

Blood Collection Tube (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	Product Description	Blood Collection Tube
	Usage	Single-Use (Disposable)
	Sterility	Sterile
TECHNICAL INFORMATION	Type of Tube	Vacuum Blood Collection Tube (Evacuated Tube)
	Material of Tube	Polyethylene Terephthalate (PET)
	Non toxic, leakproof and crack resistance material	Yes
	Transparency	Transparent
	Negative air pressure present inside the tube	Yes
	Additive	Sodium Fluoride & EDTA
	Draw volume of blood sample	2 mL
	Cap Color	Grey
	Patient Utility	Adult

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
	Graduation marking	Yes
	Label on tube for Identification	Yes
CERTIFICATIONS	Compliance to Medical Device Rules (MDR) 2017 as amended till date	Yes
	Availability of valid medical device license for the product issued from the competent authority defined under Drugs and Cosmetic Act 1940 and Rules made there under as amended till date	Yes
	Manufacturer Certification	ISO:13485 (Latest)
	Availability of Test Report for each supplied batch/product as per Medical Device Rules (MDR) 2017 as amended till date	Yes
	Submission of all necessary certifications, licenses and test reports to the buyer at the time of bid submission and/or along with supplies as per buyer requirement	Yes
SHELF LIFE	Minimum shelf life of the product at the time of delivery to the consignee	3/4 th of Total Shelf Life
ADVANCE SAMPLE	Agree to provide advance sample of the product for buyer's approval after award of contract and before commencement of supply in case of bidding only	Yes

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

क्र.सं./S.N o.	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.	प्रेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer	पता/Address	डिलीवरी अनुसूची /Delivery Schedule अनुबंध प्रारम्भ होने की तारीख से दिनों की संख्या में / (In number of days from contract start days)									
1	Rakesh Choudhary	181133,ESIC MODEL HOSPITAL, LANE NO.-02 - SIDCO, OPPOSITE POLICE STATION, BARI BRAHMANA JAMMU	<table border="1"> <thead> <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> </thead> <tbody> <tr> <td>1000</td> <td>1</td> <td>15</td> </tr> <tr> <td>1000</td> <td>150</td> <td>180</td> </tr> </tbody> </table>	मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____ तक पूरी कर ली जाए /Delivery to be completed by	1000	1	15	1000	150	180
मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____ तक पूरी कर ली जाए /Delivery to be completed by										
1000	1	15										
1000	150	180										

Blood Collection Tube (15000 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	Product Description	Blood Collection Tube
	Usage	Single-Use (Disposable)
	Sterility	Sterile
TECHNICAL INFORMATION	Type of Tube	Vacuum Blood Collection Tube (Evacuated Tube)
	Material of Tube	Polyethylene Terephthalate (PET)
	Non toxic, leakproof and crack resistance material	Yes
	Transparency	Transparent
	Negative air pressure present inside the tube	Yes
	Additive	K2 EDTA
	Draw volume of blood sample	3 mL
	Cap Color	Lavender or Purple
Patient Utility	Adult	

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
	Graduation marking	Yes
	Label on tube for Identification	Yes
CERTIFICATIONS	Compliance to Medical Device Rules (MDR) 2017 as amended till date	Yes
	Availability of valid medical device license for the product issued from the competent authority defined under Drugs and Cosmetic Act 1940 and Rules made there under as amended till date	Yes
	Manufacturer Certification	ISO:13485 (Latest)
	Availability of Test Report for each supplied batch/product as per Medical Device Rules (MDR) 2017 as amended till date	Yes
	Submission of all necessary certifications, licenses and test reports to the buyer at the time of bid submission and/or along with supplies as per buyer requirement	Yes
SHELF LIFE	Minimum shelf life of the product at the time of delivery to the consignee	3/4 th of Total Shelf Life
ADVANCE SAMPLE	Agree to provide advance sample of the product for buyer's approval after award of contract and before commencement of supply in case of bidding only	Yes

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

क्र.सं./S.N o.	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 प्रारम्भ होने की तारीख से दिनों की संख्या में / (In number of days from contract start days)												
1	Rakesh Choudhary	181133,ESIC MODEL HOSPITAL, LANE NO.-02 - SIDCO, OPPOSITE POLICE STATION, BARI BRAHMANA JAMMU	<table border="1"> <thead> <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> </thead> <tbody> <tr> <td>5000</td> <td>0</td> <td>15</td> </tr> <tr> <td>5000</td> <td>105</td> <td>135</td> </tr> <tr> <td>5000</td> <td>210</td> <td>240</td> </tr> </tbody> </table>	मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____ तक पूरी कर ली जाए /Delivery to be completed by	5000	0	15	5000	105	135	5000	210	240
मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____ तक पूरी कर ली जाए /Delivery to be completed by													
5000	0	15													
5000	105	135													
5000	210	240													

Blood Collection Tube (15000 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 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)
GENERAL	Product Description	Blood Collection Tube
	Usage	Single-Use (Disposable)
	Sterility	Sterile
TECHNICAL INFORMATION	Type of Tube	Vacuum Blood Collection Tube (Evacuated Tube)
	Material of Tube	Polyethylene Terephthalate (PET)
	Non toxic, leakproof and crack resistance material	Yes
	Transparency	Transparent
	Negative air pressure present inside the tube	Yes
	Additive	Clot Activator without Serum Separation Gel
	Draw volume of blood sample	4 mL
Cap Color	Red	

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
	Patient Utility	Adult
	Graduation marking	Yes
CERTIFICATIONS	Compliance to Medical Device Rules (MDR) 2017 as amended till date	Yes
	Availability of valid medical device license for the product issued from the competent authority defined under Drugs and Cosmetic Act 1940 and Rules made there under as amended till date	Yes
	Availability of Test Report for each supplied batch/product as per Medical Device Rules (MDR) 2017 as amended till date	Yes
	Submission of all necessary certifications, licenses and test reports to the buyer at the time of bid submission and/or along with supplies as per buyer requirement	Yes
SHELF LIFE	Minimum shelf life of the product at the time of delivery to the consignee	3/4 th of Total Shelf Life
ADVANCE SAMPLE	Agree to provide advance sample of the product for buyer's approval after award of contract and before commencement of supply in case of bidding only	Yes, No

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

क्र.सं./S.N o.	परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer	पता/Address	डिलीवरी अनुसूची /Delivery Schedule अनुबंध प्रारम्भ 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 प्रारंभ होने की तारीख से दिनों की संख्या में / (In number of days from contract start days)		
1	Rakesh Choudhary	181133,ESIC MODEL HOSPITAL, LANE NO.-02 - SIDCO, OPPOSITE POLICE STATION, BARI BRAHMANA JAMMU	मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____ तक पूरी कर ली जाए /Delivery to be completed by
			5000	1	15
			5000	105	135
			5000	210	240

Blood Collection Tube (15000 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	Product Description	Blood Collection Tube
	Usage	Single-Use (Disposable)
	Sterility	Sterile
TECHNICAL INFORMATION	Type of Tube	Vacuum Blood Collection Tube (Evacuated Tube)
	Material of Tube	Polyethylene Terephthalate (PET)
	Non toxic, leakproof and crack resistance material	Yes
	Transparency	Transparent
	Negative air pressure present inside the tube	Yes
	Additive	Clot Activator with Serum Separation Gel
	Draw volume of blood sample	4 mL
	Cap Color	Yellow/Golden
	Patient Utility	Adult
	Graduation marking	Yes
Label on tube for Identification	Yes	

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
CERTIFICATIONS	Compliance to Medical Device Rules (MDR) 2017 as amended till date	Yes
	Availability of valid medical device license for the product issued from the competent authority defined under Drugs and Cosmetic Act 1940 and Rules made there under as amended till date	Yes
	Manufacturer Certification	ISO:13485 (Latest)
	Availability of Test Report for each supplied batch/product as per Medical Device Rules (MDR) 2017 as amended till date	Yes
	Submission of all necessary certifications, licenses and test reports to the buyer at the time of bid submission and/or along with supplies as per buyer requirement	Yes
SHELF LIFE	Minimum shelf life of the product at the time of delivery to the consignee	3/4 th of Total Shelf Life
ADVANCE SAMPLE	Agree to provide advance sample of the product for buyer's approval after award of contract and before commencement of supply in case of bidding only	Yes

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

क्र.सं./S.N o.	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 प्रारम्भ होने की तारीख से दिनों की संख्या में / (In number of days from contract start days)		
1	Rakesh Choudhary	181133,ESIC MODEL HOSPITAL, LANE NO.-02 - SIDCO, OPPOSITE POLICE STATION, BARI BRAHMANA JAMMU	मात्रा /Quantity	प्रारंभ होने की तारीख से डिलीवरी /Delivery to start after	डिलीवरी _____ तक पूरी कर ली जाए /Delivery to be completed by
			5000	1	15
			5000	105	135
			5000	210	240

Blood Collection Needles (V2) (15000 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	Product Description	Blood Collection Needle
	Utility	Multi-sample draw Needle
	Usage	Single-Use
	Sterility	Sterile
DIMENSIONS	Needle Length (in mm)	25.0 (millimeter)
	Needle Size	22G
MATERIAL & CONSTRUCTION	Material of Needle	Medical Grade Stainless Steel
	Integrated Safety Shield that Fully Covers the Needle After Use	Yes
PACKAGING	Type of Packing	Individually packed in a twist apart plastic container
CERTIFICATIONS & REPORTS	Compliance to Medical Device Rules (MDR) 2017 as amended till date	Yes
	Availability of valid medical device license for the product issued from the competent authority defined under Drugs and Cosmetic Act 1940 and Rules made there under as amended till date	Yes

विवरण/Specification	विशिष्टि का नाम /Specification Name	बिड के लिए आवश्यक अनुमत मूल्य /Bid Requirement (Allowed Values)
SHELF LIFE	Minimum shelf life of the product at the time of delivery to the consignee	3/4 th of Total Shelf Life

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 प्रारम्भ 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	Rakesh Choudhary	181133,ESIC MODEL HOSPITAL, LANE NO.-02 - SIDCO, OPPOSITE POLICE STATION, BARI BRAHMANA JAMMU			
			5000	1	15
			5000	105	135
			5000	210	240

Special terms and conditions-Version:1 effective from 30-09-2025 for category Rapid Test Kit for HIV / Syphilis / Hepatitis B Virus (HBV) / Hepatitis C Virus (HCV)

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.

Special terms and conditions-Version:1 effective from 30-09-2025 for category Point of Care Rapid

Test Kits for Humans - Dengue, Malaria, Typhoid & Others

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.

Special terms and conditions-Version:1 effective from 06-10-2025 for category Rapid Pregnancy Test Kit

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.

Special terms and conditions-Version:1 effective from 01-01-2025 for category Blood Collection Tube

1. 1. All Provisions of Drugs and Cosmetics Act, 1940 and Rules (including Medical Device Rule 2017) made there under as amended 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 time to time in this regard.
2. The sellers are registered on GeM based on the self declaration of valid Medical Device 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 license, product certification, manufacturer certification/licenses, test reports etc.
3. In case of authorized resellers/distributors, it will be the legal & regulatory liability of the manufacturer to ensure that their resellers/distributors are operating in compliance with all relevant laws and regulations and are properly licensed to sell the manufacturer's products, including verifying the validity and authenticity of license held by them.

4. The price offered by the seller/bidder shall not, in any case exceed the DPCO/NPPA controlled price or price fixed by State Government, if any. The seller must reduce the prices if there is any reduction in DPCO/NPPA ceiling price or price fixed by State Government, if any.
5. 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. ATC shall supersede specific STC which shall supersede General Terms and Conditions (GTC), whenever there are any conflicting provisions.

Special terms and conditions-Version:2 effective from 27-03-2026 for category Blood Collection Needles (V2)

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.

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. 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.

3. Inspection

Testing of Sample: The testing of advance sample and bulk sample during PDI will be carried at the designated AHSP labs. When testing facilities are not available, the facilities of Govt labs/NABL/Accredited labs will be utilized. The testing charges outside the designated AHSP labs to be borne by seller's.

4. **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 10 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.

5. **Forms of EMD and PBG**

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

Medical superintendent
payable at
ESIC Fund account no 1

. 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. **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

Medical superintendent
payable at
ESIC Fund account no 1

. 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.

7. **Financial Criteria**

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

8. **Rate Contract**

AVAILABILITY OF OFFICE OF SERVICE PROVIDER: An office of the Service Provider must be located in the state of Consignee. DOCUMENTARY EVIDENCE TO BE SUBMITTED.

9. **Scope of Supply**

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

10. **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.

11. **Certificates**

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

12. **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.

13. **Certificates**

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

as per ATC

14. **Buyer Added Bid Specific ATC**

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

15. **Warranty**

Timely Servicing / rectification of defects during warranty period: After having been notified of the defects / service requirement during warranty period, Seller has to complete the required Service / Rectification within 7 days time limit. If the Seller fails to complete service / rectification with defined time limit, a penalty of 0.5% of Unit Price of the product shall be charged as penalty for each week of delay from the seller. Seller can deposit the penalty with the Buyer directly else the Buyer shall have a right to recover all such penalty amount from the Performance Security (PBG). Cumulative Penalty cannot exceed more than 10% of the total contract value after which the Buyer shall have the right to get the service / rectification done from alternate sources at the risk and cost of the Seller besides forfeiture of PBG. Seller shall be liable to reimburse the cost of such service / rectification to the Buyer.

16. **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.

17. **Buyer Added Bid Specific ATC**

Buyer Added text based ATC clauses

ADDITIONAL TERMS & CONDITIONS (ATC)

For Procurement of Rapid Diagnostic Kits & Blood Collection Consumables

ESIC Model Hospital, Bari Brahmana, Jammu

1. Scope of Supply

The bidder shall supply the following items:

A. Rapid Diagnostic Test Kits

- HCV Rapid Cards (Single Test)
- HIV Rapid Cards (Single Test)
- HBsAg Rapid Cards (Single Test)
- VDRL Rapid Cards (Single Test)

- Pregnancy Cards (Single Test)
- Malaria Cards (Single Test)
- Chikungunya Cards (Single Test)
- Widal Typhi IgG + IgM Cards (Single Test)
- Dengue Cards (NS1/IgG/IgM as applicable)
- Occult Blood Test Kits (50 Tests Pack)

B. Blood Collection Consumables

- Vacutainer Needles (22G)
- Clot Activator Tubes – 4 ml
- K2 EDTA Tubes (Purple) – 3 ml
- Plain Tubes (Red) – 4 ml
- Sodium Fluoride Tubes (Grey) – 2 ml
- Sodium Citrate 3.2% Tubes (Blue) – 1.8 ml
- Trisodium Citrate 3.8% Tubes (Black) – 2 ml

2. Mandatory Certifications (STRICT - REJECTION IF NOT COMPLIED)

For Rapid Test Kits

- Must be **DCGI Approved / Licensed**
- Must comply with **IVD standards**
- Should have **WHO-GMP / ISO 13485 certification**
- For HIV kits: **NACO approved preferred**
- Sensitivity \geq **99%**, Specificity \geq **99%** (document proof required)

For Vacutainers & Needles

- Must be **CE Certified / ISO 13485**
- Sterile, single-use, non-pyrogenic
- Vacuum tubes must be **leak-proof & pre-calibrated**

3. Technical Compliance (MANDATORY DOCUMENT)

Bidder must upload:

- **Item-wise compliance sheet** Annexure A
- Product brochure/catalogue
- Regulatory certificates (DCGI / CE / ISO)
- Test performance data (Sensitivity/Specificity)
- Annexure B (Undertaking)

Non-submission = Bid Rejection

4. Make / Brand Criteria (VERY IMPORTANT)

- Only **reputed / established brands** allowed
- No **local / unbranded / assembled products**
- Bidder must clearly mention **Brand Name & Model**

5. Shelf Life Requirement

- Minimum **75% shelf life at the time of supply**
- Kits with less shelf life \rightarrow **Rejected**

6. Sample / Demonstration Clause

- Buyer reserves right to:
 - Ask for **sample before finalization**
 - Conduct **performance validation**

- If sample fails → **Bid rejected**

7. Packaging & Storage

- Proper **temperature-controlled packaging**
- No damaged / expired / tampered supply
- Each pack must have:
 - Batch No.
 - Manufacturing & Expiry date

8. Delivery Conditions

- Staggered Delivery schedule after order within **15-30 days**
- Delivery at:
ESIC Model Hospital, Bari Brahmna, Jammu
- Delay penalty applicable as per GeM rules

9. Replacement Clause

- Defective / non-performing items must be:
 - Replaced within **7 days**
- No extra cost to buyer

10. Performance Guarantee

- Supplier must ensure:
 - Proper functioning of test kits
 - Accurate and reproducible results

11. Past Experience Criteria (QUALITY FILTER)

Bidder must submit:

- At least **2 similar supply orders**
- Preferably to:
 - Government hospitals / ESIC / AIIMS / Medical Colleges

12. Undertaking by Bidder as per Proforma attached

Bidder must submit signed declaration:

- Products are **genuine & certified**
- No **spurious / substandard material**
- All specifications are complied

13. Inspection Clause

- Buyer/committee may inspect:
 - At delivery stage
 - Or randomly during use
- Non-compliance → **Blacklisting + Rejection**

14. Payment Terms

- Payment after:
 - Successful delivery
 - Verification by user department

15. Bid Evaluation Criteria

- L1 will be decided **item-wise / overall (as per bid setup)**
- However, only **technically qualified bidders** will be considered

16. Rejection Criteria (STRICT)

Bid will be rejected if:

- No certifications uploaded
- No compliance sheet

- Low-quality / unbranded items quoted
- False documents submitted

TECHNICAL COMPLIANCE SHEET(ANNEXURE A)

Bid No.: _____

Name of Bidder: _____

Hospital: ESIC Model Hospital, Bari Brahmana, Jammu

Sr. No.	Item Name	Specification Required	Offered Specification	Make/Brand	Model	Compliance (Yes/No)	Remarks
1	HCV Rapid Cards	Single Test, DCGI Approved, Sensitivity $\geq 99\%$, Specificity $\geq 99\%$					

2	HIV Rapid Cards	Single Test, HIV 1&II NACO/DCGI Approved, High Sensitivity/Specificity					
3	HBsAg Rapid Cards	Single Test, IVD Certified, High Accuracy					
4	VDRL Rapid Cards	Single Test, Approved Kit					
5	Pregnancy Cards	Single Test, High Sensitivity					
6	Malaria Cards	Pf/Pv Detection, Rapid Kit					
7	Chikungunya Cards	IgM Detection					
8	Widal Typhi IgG + IgM	Rapid Test					
9	Dengue Cards	NS1/IgG/IgM Combo					
10	Occult Blood Test	50 Tests Pack					
11	Vacutainer Needles	22G, Sterile, Disposable					
12	Clot Activator Tube	4 ml, Sterile, Vacuum					
13	K2 EDTA Tube	3 ml, Purple Cap					
14	Plain Tube Red	4 ml					

15	Sodium Fluoride Tube	2 ml, Grey Cap					
16	Sodium Citrate 3.2%	1.8 ml, Blue Cap					
17	Trisodium Citrate 3.8%	2 ml, Black Cap					

Mandatory Declaration (to be added at bottom)

We hereby certify that all the offered products meet the required specifications, are certified (DCGI/CE/ISO), and comply with all terms & conditions of the bid.

Authorized Signatory

Name: _____
Seal & Signature

UNDERTAKING FORMAT

(ANNEXURE B)

UNDERTAKING / DECLARATION BY BIDDER

Bid No.: _____

Date: _____

To,
The Medical Superintendent
ESIC Model Hospital
Bari Brahmana, Jammu

Subject: Undertaking for Supply of Rapid Test Kits & Blood Collection Consumables

I/We, **(Name of Bidder)**, hereby undertake the following:

1. ✓ All products supplied shall be **new, unused, and of standard quality**.
2. ✓ All diagnostic kits are:
 - **DCGI approved / Licensed**
 - Compliant with **IVD standards**
 - Having **Sensitivity & Specificity ≥ 99%** (where applicable)
3. ✓ All vacutainers and needles are:
 - **Sterile, single-use**
 - **CE / ISO 13485 certified**
4. ✓ No **spurious, substandard, or duplicate products** will be supplied.
5. ✓ Minimum **75% shelf life** will be available at the time of delivery.
6. ✓ We agree to provide **sample/testing** if demanded by the buyer.
7. ✓ Any defective or non-performing item will be:
 - Replaced within **7 days**
 - At **no extra cost**
8. ✓ We have **adequate experience** in supplying similar items to Government institutions.
9. ✓ All documents submitted are **true and genuine**.
In case of false information:
 - Buyer may **cancel bid**

- Initiate **blacklisting**
10. ✓ We agree to all **ATC conditions** mentioned in the bid document.
-

Declaration

I/We certify that the above information is true to the best of my knowledge and belief.

Authorized Signatory

Name: _____
Designation: _____
Firm Name: _____
Seal & Signature

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 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---