बिड संख्या/Bid Number: GEM/2026/B/7449153

दिनांक /Dated: 27-04-2026

बिड दस्तावेज़ / Bid Document

बिड विवरण/Bid Details	
बिड बंद होने की तारीख/समय /Bid End Date/Time	18-05-2026 11:00:00
बिड खुलने की तारीख/समय /Bid Opening Date/Time	18-05-2026 11:30:00
बिड पेशकश वैधता (बंद होने की तारीख से)/Bid Offer Validity (From End Date)	90 (Days)
मंत्रालय/राज्य का नाम/Ministry/State Name	Pmo
विभाग का नाम/Department Name	Department Of Atomic Energy
संगठन का नाम/Organisation Name	Nuclear Power Corporation Of India Limited
कार्यालय का नाम/Office Name	Naps
कुल मात्रा/Total Quantity	5600
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) , Bivalent RDT Malaria Kit for NCVBDC Programme (Q1) , Widal Test Kit - Agglutination Method for Human Samples (Q2) , Rheumatoid Factor Test Kit - RA Test Kit (V2) (Q2) , ASO (Anti Streptolysin-O) Test Kits (V2) (Q2)
वर्षों के अनुभव एवं टर्नओवर से एमएसई को छूट प्राप्त है / MSE Relaxation for Years Of Experience and Turnover	Yes | Complete
स्टार्टअप के लिए अनुभव के वर्षों और टर्नओवर से छूट प्रदान की गई है / Startup Relaxation for Years Of Experience and Turnover	Yes | Complete
बिड लगाने की समय सीमा स्वतः नहीं बढ़ाने के लिए आवश्यक बिड की संख्या। / Minimum number of bids required to disable automatic bid extension	3
दिनों की संख्या, जिनके लिए बिड लगाने की समय-सीमा बढ़ाई जाएगी। / Number of days for which Bid would be auto-extended	7
ऑटो एक्सटेंशन अधिकतम कितनी बार किया जाना है। / Number of Auto Extension count	1
बिड से रिवर्स नीलामी सक्रिय किया/Bid to RA enabled	No
बिड का प्रकार/Type of Bid	Two Packet Bid

बिड विवरण/Bid Details	
प्राथमिक उत्पाद श्रेणी/Primary product category	Rapid Test Kit for HIV / Syphilis / Hepatitis B Virus (HBV) / Hepatitis C Virus (HCV)
तकनीकी मूल्यांकन के दौरान तकनीकी स्पष्टीकरण हेतु अनुमत समय /Time allowed for Technical Clarifications during technical evaluation	7 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
Payment Timelines	Payments shall be made to the Seller within **10** days of issue of consignee receipt-cum-acceptance certificate (CRAC) and on-line submission of bills (This is in supersession of 10 days time as provided in clause 12 of GeM GTC)
मूल्यांकन पद्धति/Evaluation Method	Total value wise evaluation
मध्यस्थता खंड/Arbitration Clause	No
सुलह खंड/Mediation Clause	No

ईएमडी विवरण/EMD Detail

आवश्यकता/Required	No

ईपीबीजी विवरण /ePBG Detail

आवश्यकता/Required	No

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

एमआईआई खरीद वरीयता/MII Purchase Preference

एमआईआई खरीद वरीयता/MII Purchase Preference	No

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

ट्रेड्स भुगतान संबंधी विवरण/TReDS Payment Details

This Bid provides for Trade Receivables Discounting System (TReDS) as Preferred mode of payment. For MSME sellers, payments may be processed through a TReDS exchange in which the Buyer is registered, subject to applicable policy and regulatory guidelines. Accordingly, sellers intending to avail payment through TReDS are required to be registered with at least one TReDS exchange in which the buyer is registered.

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. 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.
6. Estimated Bid Value indicated above is being declared solely for the purpose of guidance on EMD amount and for determining the Eligibility Criteria related to Turn Over, Past Performance and Project / Past Experience etc. This has no relevance or bearing on the price to be quoted by the bidders and is also not going to have any impact on bid participation. Also this is not going to be used as a criteria in determining reasonableness of quoted prices which would be determined by the buyer based on its own assessment of reasonableness and based on competitive prices received in Bid / RA process.

एक्सेल में अपलोड किए जाने की आवश्यकता /Excel Upload Required :

GST BREAKUP TO BE UPLOAED IN FINANCIAL BID OR PRICE BID ONLY - 1776403004.xlsx

Rapid Test Kit For HIV / Syphilis / Hepatitis B Virus (HBV) / Hepatitis C Virus (HCV) (500 packet)

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
	Result Time	? 30 minutes
	Positive and negative controls provided with each pack of kit	No
PACKAGING	**Number of Tests per Pack**	25 Tests
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	मात्रा /Quantity	डिलीवरी के दिन/Delivery Days
1	Ram Resh	202389,Manager (Materials),-C&MM, Narora Atomic Power Station, Distt.-Bulandshahar (UP), PIN – 203389	500	30

Point Of Care Rapid Test Kits For Humans – Dengue, Malaria, Typhoid & Others (500 packet)

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
	Result Time	? 15 minutes
	Positive and negative controls provided with each pack of kit	No
PACKAGING	**Number of Tests per Pack**	10 Tests
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	मात्रा /Quantity	डिलीवरी के दिन/Delivery Days
1	Ram Resh	202389,Manager (Materials),-C&MM, Narora Atomic Power Station, Distt.-Bulandshahar (UP), PIN – 203389	500	30

Point Of Care Rapid Test Kits For Humans – Dengue, Malaria, Typhoid & Others (500 packet)

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
	Result Time	? 15 minutes
	Positive and negative controls provided with each pack of kit	No
PACKAGING	**Number of Tests per Pack**	10 Tests

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

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

क्र.सं./S.No.	परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer	पता/Address	मात्रा /Quantity	डिलीवरी के दिन/Delivery Days
1	Ram Resh	202389,Manager (Materials),-C&MM, Narora Atomic Power Station, Distt.-Bulandshahar (UP), PIN – 203389	500	30

Bivalent RDT Malaria Kit For NCVBDC Programme (500 Test)

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	**Conformity to technical specifications**	As per detailed technical specification document attached/uploaded

अतिरिक्त विशिष्टि दस्तावेज़ /Additional Specification Documents

Applicable Specification Document	View

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

क्र.सं./S.No.	परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer	पता/Address	मात्रा /Quantity	डिलीवरी के दिन/Delivery Days
1	Ram Resh	202389,Manager (Materials),-C&MM, Narora Atomic Power Station, Distt.-Bulandshahar (UP), PIN – 203389	500	30

Widal Test Kit - Agglutination Method For Human Samples (3000 packet)

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**	Widal Test Kit (Slide Test)
	Type of antigen set provided	S. paratyphi 'AH', S. paratyphi 'BH', S. typhi 'H, S. typhi 'O'
	Result Type	Qualitative
	Sample Type	Serum
	Positive and Negative control provided	Yes
PACKAGING	**Total Volume of Reagent per Pack**	20 ml
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	मात्रा /Quantity	डिलीवरी के दिन/Delivery Days
1	Ram Resh	202389,Manager (Materials),-C&MM, Narora Atomic Power Station, Distt.-Bulandshahar (UP), PIN – 203389	3000	30

Rheumatoid Factor Test Kit - RA Test Kit (V2) (300 Test)

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 Description	**Testing Principle**	Latex Agglutination Slide Method
Packaging	**Pack Size**	50 Tests

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

क्र.सं./S.No.	परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer	पता/Address	मात्रा /Quantity	डिलीवरी के दिन/Delivery Days
1	Ram Resh	202389,Manager (Materials),-C&MM, Narora Atomic Power Station, Distt.-Bulandshahar (UP), PIN – 203389	300	30

ASO (Anti Streptolysin-O) Test Kits (V2) (300 Test)

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 Informations	**Result Type**	Qualitative
	Sensitivity	? 200 IU/ml for Agglutination Slide Method
Packaging	**Pack size of kit**	100 Tests

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

क्र.सं./S.No.	परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer	पता/Address	मात्रा /Quantity	डिलीवरी के दिन/Delivery Days
1	Ram Resh	202389,Manager (Materials),-C&MM, Narora Atomic Power Station, Distt.-Bulandshahar (UP), PIN – 203389	300	30

क्र.सं./S.No.	परेषिती/रिपोर्टिंग अधिकारी /Consignee Reporting/Officer	पता/Address	मात्रा /Quantity	डिलीवरी के दिन/Delivery Days

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. 1. *A*ll Provisions of Drugs and Cosmetics Act, 1940 and Rules (including Medical Device Rule 2017) made there under and all subsequent amendments till date will always be applicable. This will include all notifications issued by Central Drugs Standard Control Organization (CDSCO), Ministry of Health & Family Welfare (MoHFW) and Department of Pharmaceuticals (DOP), Ministry of Chemicals & Fertilizers in this regard.
   2. The seller’s registration and product upload on GeM is based on their declaration of possessing a valid Medical Device License for the product, certifications, test reports as per MDR (2017), latest amended. However, buyers must check, validate and verify the details at their end for all applicable licenses and certifications e.g., validity and authenticity/genuineness of medical device license (or registration), product certifications, manufacturer’s certifications/ licenses, test reports at the time of supply.
   3. In case of authorized resellers/distributors, it will be the legal and regulatory liability of the manufacturer to ensure that their resellers/distributors are operating in compliance with all relevant laws and regulations, and are licensed to sell the manufacturer's products, including verifying the validity and authenticity of license held by them.
   4. Any other Terms and Conditions which is not included or at variance with the conditions specified in STC/GTC, may be added by the buyer through Additional Terms and Conditions (ATC) in the bid to ensure items are procured from authentic/validated source with appropriate and applicable quality. The above terms and conditions are in reverse order of precedence i.e. BID ATC shall supersede specific STC which shall supersede General Terms and Conditions (GTC), whenever there are any conflicting provisions.
   5. **Packing and Marking:** Should be as per MDR.

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. *A*ll Provisions of Drugs and Cosmetics Act, 1940 and Rules (including Medical Device Rule 2017) made there under and all subsequent amendments till date will always be applicable. This will include all notifications issued by Central Drugs Standard Control Organization (CDSCO), Ministry of Health & Family Welfare (MoHFW) and Department of Pharmaceuticals (DOP), Ministry of Chemicals & Fertilizers in this regard.
   2. The seller’s registration and product upload on GeM is based on their declaration of possessing a valid Medical Device License for the product, certifications, test reports as per MDR (2017), latest amended. However, buyers must check, validate and verify the details at their end for all applicable licenses and certifications e.g., validity and authenticity/genuineness of medical device license (or registration), product certifications, manufacturer’s certifications/ licenses, test reports at the time of supply.
   3. In case of authorized resellers/distributors, it will be the legal and regulatory liability of the manufacturer to ensure that their resellers/distributors are operating in compliance with all relevant laws and regulations, and are licensed to sell the manufacturer's products, including verifying the validity and authenticity of license held by them.
   4. Any other Terms and Conditions which is not included or at variance with the conditions specified in STC/GTC, may be added by the buyer through Additional Terms and Conditions (ATC) in the bid to ensure items are procured from authentic/validated source with appropriate and applicable quality. The above terms and conditions are in reverse order of precedence i.e. BID ATC shall supersede specific STC which shall supersede General Terms and Conditions (GTC), whenever there are any conflicting provisions.
   5. **Packing and Marking:** Should be as per MDR.

Special terms and conditions-Version:2 effective from 27-03-2026 for category Bivalent RDT Malaria

Kit for NCVBDC Programme

1. 1. *A*ll Provisions of Drugs and Cosmetics Act, 1940 and Rules (including Medical Device Rule 2017) made there under and all subsequent amendments till date will always be applicable. This will include all notifications issued by Central Drugs Standard Control Organization (CDSCO), Ministry of Health & Family Welfare (MoHFW) and Department of Pharmaceuticals (DOP), Ministry of Chemicals & Fertilizers in this regard.
   2. The seller’s registration and product upload on GeM is based on their declaration of possessing a valid Medical Device License for the product, certifications, test reports as per MDR (2017), latest amended. However, buyers must check, validate and verify the details at their end for all applicable licenses and certifications e.g., validity and authenticity/genuineness of medical device license (or registration), product certifications, manufacturer’s certifications/ licenses, test reports at the time of supply.
   3. In case of authorized resellers/distributors, it will be the legal and regulatory liability of the manufacturer to ensure that their resellers/distributors are operating in compliance with all relevant laws and regulations, and are licensed to sell the manufacturer's products, including verifying the validity and authenticity of license held by them.
   4. Any other Terms and Conditions which is not included or at variance with the conditions specified in STC/GTC, may be added by the buyer through Additional Terms and Conditions (ATC) in the bid to ensure items are procured from authentic/validated source with appropriate and applicable quality. The above terms and conditions are in reverse order of precedence i.e. BID ATC shall supersede specific STC which shall supersede General Terms and Conditions (GTC), whenever there are any conflicting provisions.
   5. **Packing and Marking:**Should be as per MDR.

Special terms and conditions-Version:1 effective from 30-10-2025 for category Widal Test Kit - Agglutination Method for Human Samples

1. 1. *A*ll Provisions of Drugs and Cosmetics Act, 1940 and Rules (including Medical Device Rule 2017) made there under and all subsequent amendments till date will always be applicable. This will include all notifications issued by Central Drugs Standard Control Organization (CDSCO), Ministry of Health & Family Welfare (MoHFW) and Department of Pharmaceuticals (DOP), Ministry of Chemicals & Fertilizers in this regard.
   2. The seller’s registration and product upload on GeM is based on their declaration of possessing a valid Medical Device License for the product, certifications, test reports as per MDR (2017), latest amended. However, buyers must check, validate and verify the details at their end for all applicable licenses and certifications e.g., validity and authenticity/genuineness of medical device license (or registration), product certifications, manufacturer’s certifications/ licenses, test reports at the time of supply.
   3. In case of authorized resellers/distributors, it will be the legal and regulatory liability of the manufacturer to ensure that their resellers/distributors are operating in compliance with all relevant laws and regulations, and are licensed to sell the manufacturer's products, including verifying the validity and authenticity of license held by them.
   4. Any other Terms and Conditions which is not included or at variance with the conditions specified in STC/GTC, may be added by the buyer through Additional Terms and Conditions (ATC) in the bid to ensure items are procured from authentic/validated source with appropriate and applicable quality. The above terms and conditions are in reverse order of precedence i.e. BID ATC shall supersede specific STC which shall supersede General Terms and Conditions (GTC), whenever there are any conflicting provisions.
   5. **Packing and Marking:** Should be as per MDR.

Special terms and conditions-Version:2 effective from 10-02-2026 for category Rheumatoid Factor Test Kit - RA Test Kit (V2)

1. 1. *A*ll Provisions of Drugs and Cosmetics Act, 1940 and Rules (including Medical Device Rule 2017) made there under and all subsequent amendments till date will always be applicable. This will include all notifications issued by Central Drugs Standard Control Organization (CDSCO), Ministry of Health & Family Welfare (MoHFW) and Department of Pharmaceuticals (DOP), Ministry of Chemicals & Fertilizers in this regard.
   2. The seller’s registration and product upload on GeM is based on their declaration of possessing a valid Medical Device License for the product, certifications, test reports as per MDR (2017), latest amended. However, buyers must check, validate and verify the details at their end for all applicable licenses and certifications e.g., validity and authenticity/genuineness of medical device license (or registration), product certifications, manufacturer’s certifications/ licenses, test reports at the time

of supply.
3. In case of authorized resellers/distributors, it will be the legal and regulatory liability of the manufacturer to ensure that their resellers/distributors are operating in compliance with all relevant laws and regulations, and are licensed to sell the manufacturer's products, including verifying the validity and authenticity of license held by them.
4. Any other Terms and Conditions which is not included or at variance with the conditions specified in STC/GTC, may be added by the buyer through Additional Terms and Conditions (ATC) in the bid to ensure items are procured from authentic/validated source with appropriate and applicable quality. The above terms and conditions are in reverse order of precedence i.e. BID ATC shall supersede specific STC which shall supersede General Terms and Conditions (GTC), whenever there are any conflicting provisions.
5. **Packing and Marking:** Should be as per MDR.

Special terms and conditions-Version:2 effective from 25-03-2026 for category ASO (Anti Streptolysin-O) Test Kits (V2)

1. 1. *A*ll Provisions of Drugs and Cosmetics Act, 1940 and Rules (including Medical Device Rule 2017) made there under and all subsequent amendments till date will always be applicable. This will include all notifications issued by Central Drugs Standard Control Organization (CDSCO), Ministry of Health & Family Welfare (MoHFW) and Department of Pharmaceuticals (DOP), Ministry of Chemicals & Fertilizers in this regard.
   2. The seller’s registration and product upload on GeM is based on their declaration of possessing a valid Medical Device License for the product, certifications, test reports as per MDR (2017), latest amended. However, buyers must check, validate and verify the details at their end for all applicable licenses and certifications e.g., validity and authenticity/genuineness of medical device license (or registration), product certifications, manufacturer’s certifications/ licenses, test reports at the time of supply.
   3. In case of authorized resellers/distributors, it will be the legal and regulatory liability of the manufacturer to ensure that their resellers/distributors are operating in compliance with all relevant laws and regulations, and are licensed to sell the manufacturer's products, including verifying the validity and authenticity of license held by them.
   4. Any other Terms and Conditions which is not included or at variance with the conditions specified in STC/GTC, may be added by the buyer through Additional Terms and Conditions (ATC) in the bid to ensure items are procured from authentic/validated source with appropriate and applicable quality. The above terms and conditions are in reverse order of precedence i.e. BID ATC shall supersede specific STC which shall supersede General Terms and Conditions (GTC), whenever there are any conflicting provisions.
   5. **Packing and Marking:** Should be as per MDR.

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 (Increased quantity ÷ Original quantity) × Original delivery period (in days), subject to minimum of 30 days. If the original delivery period is less than 30 days, the additional time equals the original delivery period. The Purchaser may extend this calculated delivery duration up to the original delivery period while exercising the option clause. Bidders must comply with these terms.

2. **Scope of Supply**

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

3. **Purchase Preference (Centre)**

   Purchase preference to Micro and Small Enterprises (MSEs): Purchase preference will be given to MSEs as defined in Public Procurement Policy for Micro and Small Enterprises (MSEs) Order, 2012 dated

23.03.2012 issued by Ministry of Micro, Small and Medium Enterprises and its subsequent Orders/Notifications issued by concerned Ministry. If the bidder wants to avail the Purchase preference, the bidder must be the manufacturer of the offered product in case of bid for supply of goods. Traders are excluded from the purview of Public Procurement Policy for Micro and Small Enterprises. In respect of bid for Services, the bidder must be the Service provider of the offered Service. Relevant documentary evidence in this regard shall be uploaded along with the bid in respect of the offered product or service. If L-1 is not an MSE and MSE Seller (s) has/have quoted price within L-1+ 15% of margin of purchase preference /price band defined in relevant policy, such Seller shall be given opportunity to match L-1 price and contract will be awarded for percentage of 25% of total value.

4. **Buyer Added Bid Specific ATC**

Buyer Added text based ATC clauses

NOTES:-

1.Clarifications/Documents submitted during the representation time shall not be considered for evaluation.

2. Only manufacturers quoting for goods/ product manufactured by them shall be eligible for availing benefits under the Public Procurement Policy for MSEs Order 2012. Dealers/ Distributors/ Sole Agents/ Resellers/ Traders/ Stockiest will not be considered for benefits under the subject policy and are required to pay requisite EMD as stipulated in the bid document. Seller declaration stating that the offered product is manufactured by them shall be submitted along with bid.

3. **TReDs**

MSME Ministry, GoI vide notification dtd 02.11.2018 has mandated all CPSE to get on boarded on the TReDS. As per the above directives of MSME Ministry, NPCIL is already registered with RXIL. TReDS is an electronic platform for facilitating the financing / discounting of trade receivables of Micro, Small and Medium Enterprises (MSMEs) through multiple financiers. These receivables can be due from corporates and other buyers, including Government Departments and Public Sector Undertakings (PSUs).Both Buyer and the Seller must be registered on the TReDS platform for financing / factoring of trade receivables of the MSME Seller. Currently, there are four RBI-approved platforms: Receivables Exchange of India Ltd. (RXIL), A Treds Ltd. (Invoicemart), C2treds and Mynd Solutions Pvt. Ltd. (M1xchange).NPCIL also encourages its seller to register on RXIL trades Platform. In view of the above, you are requested to register on RXIL trades Platform at the earliest. For registration process, bill discounting & other complete details/FAQs and correctness of above information including current interest rates, terms & condition etc., seller may visit the https://www.rxil.in

4. LIQUIDATED DAMAGES as stipulated under Clause No.15 (iii) of GEM GTC shall not be applicable for the contract awarded against this bid.

5**.Documents related to release of payment**:-

For release of payment as per terms & conditions of GeM contract, supplier has to submit following documents along with supply to consignee and also upload the documents on GeM portal.

a) Original invoice as per applicable GST provisions.
b) **Annexure-F** i.e. GST declaration (format enclosed in GeM bid).
c) Seller bank details.

As payment is to be released in a time bound manner hence, non-receipt of these d ocument along with supply may lead to rejection of supplies due to non-compliance of document submission.

6. Price Bid Evaluation and Mismatch between “total price quoted by bidder in GeM Financi al bid and Total of Price of all item uploaded “Price bid breakup/SOQR” shall be governed a s per Annexure-I (Uploaded in buyer added specific ATC)

8.Banning of business dealings by NPCIL/Buyer shall be governed as per Annexure-II (Uploa ded in buyer added specific ATC).

9. Bidder is required to submit Undertaking as per**Annexure-III**(for One Bid Per Bidder Upl oaded in buyer added specific ATC).

10. **Instructions to the supplier**

A. Supplier may ensure the driver to reach plant premise with all original documents list ed below:-

a) Original Aadhaar card of driver and helpers.
b) Original Driving license of driver (for heavy duty vehicle, driver may come with he avy duty license)
c) Original RC (Registration card) of vehicle.
d) Original Fitness Certificate.
e) Road Tax must be paid till the date.
f) Original Insurance papers of vehicle.

Driver if failed to show all these documents in original state will itself responsible for No entry of vehicle & also no further communication will be made from our end.

B. The driver is requested to reach plant site from 09.00 AM to 03.00 PM of any working day.

C. Supplier may intimate the dispatch of material to the consignee with all the details of driver and helpers, one or more days before for the Gate pass process and other necess ary actions.

11. GST BREAKUP TO BE UPLOAED IN FINANCIAL BID OR PRICE BID ONLY

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5. **Buyer Added Bid Specific ATC**

Buyer uploaded ATC document Click here to view the file.

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---