

## अनुबंध | Contract



अनुबंध क्रमांक | Contract No: GEMC-511687782295358

अनुबंध तिथि | Generated Date : 30-Mar-2024

संगठन विवरण   Organisation Details	खरीदार विवरण   Buyer Details
प्ररूप   Type : State Government मंत्रालय   Ministry : - विभाग   Department : Health and Family Welfare Department Jammu and Kashmir संगठन का नाम   Organisation Name : Government Medical College and Associated Hospitals कार्यालय क्षेत्र   Office Zone: Cheif Medical Officer Udhampur	पद   Designation : DISTRICT HEALTH OFFICER UDHAMPUR संपर्क नंबर   Contact No. : 01992-270207- ईमेल आईडी   Email ID : dho-udhampur@jk.gov.in जीएसटीआईएन   GSTIN : 01ABMPD0422A1D3 पता   Address : Opposite Government Polytechnic Udhampur, UDHAMPUR, JAMMU & KASHMIR-182101, India

वित्तीय स्वीकृति विवरण   Financial Approval Detail	भुगतान प्राधिकरण विवरण   Paying Authority Details
आईएफडी सहमति   IFD Concurrence : No प्रशासनिक अनुमोदन का पदनाम   Designation of Administrative Approval: Chief Medical Officer, udhampur वित्तीय अनुमोदन का पदनाम   Designation of Financial Approval : Chief Medical Officer, udhampur	Role: PAO भुगतान का तरीका   Payment Mode: Offline पद   Designation : DISTRICT ACCOUNTS MANAGER ईमेल आईडी   Email ID : dam-udhampur@jk.gov.in जीएसटीआईएन   GSTIN : - पता   Address: OFFICE OF THE CHIEF MEDICAL OFFICER, OPPOSITE GOVT. POLYTECHNIC COLLEGE, UDHAMPUR., Udhampur, JAMMU & KASHMIR-182101, India

विक्रेता विवरण   Seller Details
जेम विक्रेता आईडी   GeM Seller ID : A0FA190000857066 कंपनी का नाम   Company Name : Jammu scientific Agencies संपर्क नंबर   Contact No. : 09697629052 ईमेल आईडी   Email ID : jammusciencegcm@gmail.com पता   Address : PLOT NO. 163, NEAR GOVT. MEDICAL COLLEGE, MAHESH PURA, BAKSHI NAGAR, Jammu, JAMMU & KASHMIR-180001, - एमएसएमई पंजीकरण संख्या   MSME Registration number : UDYAM-JK-07-0004034 एमएसई सामाजिक श्रेणी   MSE Social Category : General एमएसई लिंग श्रेणी   MSE Gender : Male जीएसटीआईएन   GSTIN: 01BXJPS1194Q1ZW

\*जिसके नाम के पक्ष में GST/TAX इनवॉइस पेश किया जाएगा | GST / Tax invoice to be raised in the name of - Buyer

वितरण निर्देश | Delivery Instructions : NA

उत्पाद विवरण   Product Details						
#	आइटम विवरण   Item Description	आइटम विवरण   Ordered Quantity	इकाई   Unit	इकाई मूल्य (INR)   Unit Price (INR)	कर विभाजन (INR)   Tax Bifurcation (INR)	मूल्य (INR में सभी शुल्क और कर सहित)   Price (Inclusive of all Duties and Taxes in INR)
1	उत्पाद का नाम   Product Name : BenesPhera HIV1 & HIV2 Antibodies Detection Rapid Test Kit ब्रांड   Brand : BenesPhera ब्रांड प्रकार   Brand Type : Registered Brand कैटलॉग की स्थिति   Catalogue Status: OEM verified catalogue कैसे बेचा जा रहा है   Selling As : OEM verified Reseller श्रेणी का नाम और चतुर्थांश   Category Name & Quadrant : HIV Rapid Test Kits (Q2) मॉडल   Model: BENP15001535 एचएसएन कोड   HSN Code: HSN not specified by seller	100	Test	42	NA	4,200
कुल ऑर्डर मूल्य   Total Order Value (in INR)						4,200

परोक्षिती विवरण   Consignee Detail						
क्र.सं.   S.No	परोक्षिती   Consignee	वस्तु   Item	लॉट नंबर   Lot No.	मात्रा   Quantity	दिनांक के बाद डिलीवरी शुरू करना है   Delivery Start After	वितरण पूरा कब तक करना है   Delivery To Be Completed By
	पद   Designation : DISTRICT HEALTH OFFICER UDHAMPUR ईमेल आईडी   Email ID : dho-udhampur@jk.gov.in संपर्क   Contact : 01992-270207-	BenesPhera HIV1 &				

1	जीएसटीआईएन   GSTIN : 01ABMPD0422A1D3 पता   Address : Opposite Government Polytechnic Udampur, UDHAMPUR, JAMMU & KASHMIR-182101, India	HIV2 Antibodies Detection Rapid Test Kit	-	100	30-Mar-2024	14-Apr-2024
<b>Product Specification for BenesPhera HIV1 &amp; HIV2 Antibodies Detection Rapid Test Kit</b>						
<b>विनिर्देश   Specification</b>	<b>उप-विनिर्देश   Sub-Spec</b>	<b>मूल्य   Value</b>				
GENERAL FEATURES	Product Description	HIV Rapid Test Kit				
	Clinical Purpose	To provide diagnosis of HIV infection				
PRODUCT INFORMATION	Type of Kit	HIV1 & HIV2 Antibodies Detection Rapid Test Kit				
	Assay Coating	Solid phase coated HIV1 and HIV2 recombinant and/or synthetic peptide antigens				
	Detection Type	Qualitative				
	Testing Principle	Lateral Flow Immunochromatographic Assay				
	Species Reactivity	Human				
	Type of Sample	Whole Blood, Serum, Plasma				
	Time to Result	≤ 30 minutes				
	Ability to evaluate negative or positive test result	Yes				
	Sensitivity	100%				
	Specificity	≥98%				
	Declared sensitivity and specificity shall be claimed by the manufacturer in the kit literature	Yes				
	Contains an internal control dot/band for the confirmation that the test has been performed correctly	Yes				
	The control dot/band able to detect the presence of human immunoglobulin and not be just a procedural control or meant merely for checking the flow of reagents or integrity of antigens except in lateral flow technology	Yes				
	Maintenance of cold chain by the supplier during storage and transportation of Kits at 2°C to 8°C and placement of cumulative time temperature indicator technology on every pack of kits	Yes				
KIT CONTENTS	Kit Contents	Test Card/Cassette with Desiccant, Sample Dropper, Assay Buffer (if any)				
	All the components shall be in the quantity as per pack size	Yes				
	Document detailing principle, component, antigen detail for antibody detection of HIV1&2, biosafety, methodologies, validity criteria, result interpretation, performance characteristic, assay limitation, mfg, exp date, storage condition provided	Yes				
	HIV positive and negative serum controls provided with the kit	Yes				
	Quantity of positive and negative controls provided	Sufficient for conducting 20% of the tests (10% negative and 10% positive controls)				
PACKAGING	Pack size of kit	50 Tests				
	The test kit packed in such a way that there is provision to conduct single test at a time	Yes				
	Each test card/cassette with desiccant individually packed in a hermetically sealed and non-permeable pouch	Yes				
CERTIFICATIONS & REPORTS	Compliance to Medical Device Rules (MDR) 2017 as amended till date	Yes				
	Availability of valid drug 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				
	Valid Drug License Number	MFG/IVD/2020/000002				
	Manufacturing unit certification	ISO:13485 (Latest)				
	Additional voluntary certification available	MFG/IVD/2020/000002				
	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 or along with supplies as per buyer requirement	Yes				
SHELF LIFE	Shelf life in months from the date of manufacture	24 month				
	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 before commencement of supply in case of bidding	Yes				
ADDITIONAL						

REQUIREMENT	Additional Requirement	NA
<b>ईपीबीजी विवरण   ePBG Detail</b>		
NA		
<b>नियम और शर्तें   Terms and Conditions</b>		
1. Special terms and conditions- Version:2 effective from 19-05-2023		
1.1		
<ol style="list-style-type: none"><li>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 &amp; Family Welfare (MoHFW) and Department of Pharmaceuticals (DOP), Ministry of Chemicals &amp; Fertilizers time to time in this regard.</li><li>2. The sellers are registered on GeM based on self-declaration of valid Drug License, product certification, test reports etc. However, buyers must check and validate the details at their end for all applicable licenses and certifications e.g., validity and authenticity/genuineness of drug license, product certification, manufacturer certification/licenses, test reports etc.</li><li>3. In case of authorized resellers/distributors, it will be the responsibility 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 drug license held by them. All legal &amp; regulatory liability in respect of the offered/supplied product shall be totally of both manufacturer and reseller/distributor.</li><li>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.</li><li>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.</li></ol>		
2. General Terms and Conditions-		
2.1 This contract is governed by the <a href="#">General Terms and Conditions</a> , conditions stipulated to this Product/Service as provided in the Marketplace.		
2.2 This Contract between the Seller and the Buyer, is for the supply of the Goods and/ or Services, detailed in the schedule above, in accordance with the General Terms and Conditions (GTC) unless otherwise superseded by Goods / Services specific Special Terms and Conditions (STC) and/ or BID/Reverse Auction Additional Terms and Conditions (ATC), as applicable		
नोट: यह सिस्टम जनरेटेड फाइल है। कोई हस्ताक्षर की आवश्यकता नहीं है। इस दस्तावेज़ का प्रिंट आउट भुगतान/लेनदेन उद्देश्य के लिए मान्य नहीं है।		
Note: This is system generated file. No signature is required. Print out of this document is not valid for payment/ transaction purpose.		