

## अनुबंध | Contract



अनुबंध क्रमांक | Contract No: GEMC-511687706512336

अनुबंध तिथि | Generated Date : 12-Mar-2024

संगठन विवरण   Organisation Details		खरीदार विवरण   Buyer Details	
प्ररूप   Type :	State Government	पद   Designation :	ACMO
मंत्रालय   Ministry :	-	संपर्क नंबर   Contact No. :	05176-272388-
विभाग   Department :	Medical Health and Family Welfare Department Uttar Pradesh	ईमेल आईडी   Email ID :	buycon528.mhfwdup.up@gembuyer.in
संगठन का नाम   Organisation Name :	N/A	जीएसटीआईएन   GSTIN :	-
कार्यालय क्षेत्र   Office Zone:	C.M.O Lalitpur	पता   Address :	C/O Office of The Cheif Medical officer, Civil line station Road, LALITPUR, UTTAR PRADESH-284403, India

वित्तीय स्वीकृति विवरण   Financial Approval Detail		भुगतान प्राधिकरण विवरण   Paying Authority Details	
आईएफडी सहमति   IFD Concurrence :	No	Role:	BUYER
प्रशासनिक अनुमोदन का पदनाम	cmo lalitpur	भुगतान का तरीका	Offline
Designation of Administrative Approval:		Payment Mode:	
वित्तीय अनुमोदन का पदनाम	cmo lalitpur	पद   Designation :	ACMO
Designation of Financial Approval :		ईमेल आईडी   Email ID :	buycon528.mhfwdup.up@gembuyer.in
		जीएसटीआईएन   GSTIN :	-
		पता   Address:	C/O Office of The Cheif Medical officer, Civil line station Road, Lalitpur, UTTAR PRADESH-284403, India

विक्रेता विवरण   Seller Details	
जेम विक्रेता आईडी   GeM Seller ID :	2QGJ210001933015
कंपनी का नाम   Company Name :	AARISHA ENTERPRISES
संपर्क नंबर   Contact No. :	08318275515
ईमेल आईडी   Email ID :	ekta275515@gmail.com
पता   Address :	175,TALAIYA,MINERVA CHAURAH,JHANSI, Jhansi, UTTAR PRADESH-284002, -
एमएसएमई पंजीकरण संख्या   MSME Registration number :	-
जीएसटीआईएन   GSTIN:	09AWIPJ3666K1Z0

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

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 : BIOGENIX (BSURE)--BIOGENIX INC. PRIVATE LIMITED HIV1 & HIV2 Antibodies Detection Rapid Test Kit ब्रांड   Brand : BIOGENIX (BSURE)--BIOGENIX INC. PRIVATE LIMITED ब्रांड प्रकार   Brand Type : Unregistered Brand कैटलॉग की स्थिति   Catalogue Status: OEM verified catalogue कैसे बेचा जा रहा है   Selling As : OEM verified Reseller श्रेणी का नाम और चतुर्भूति   Category Name & Quadrant : HIV Rapid Test Kits (Q2) मॉडल   Model: Biogenix BSURE एचएसएन कोड   HSN Code: HSN not specified by seller	1,500	Test	60	NA	90,000
कुल ऑर्डर मूल्य   Total Order Value (in INR)						90,000

परेषिती विवरण | Consignee Detail

क्र.सं.   S.No	परेषिती   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
1	पद   Designation : pharmacist ईमेल आईडी   Email ID : con405.mhfwdup.up@gembuyer.in संपर्क   Contact : 5176-274371-333333 जीएसटीआईएन   GSTIN : -	BIOGENIX (BSURE)--BIOGENIX INC. PRIVATE LIMITED HIV1 & HIV2	-	1,500	12-Mar-2024	27-Mar-2024

पता Address : C/O Office of The Cheif Medical officer, Civil line station Road, LALITPUR, UTTAR PRADESH-284403, India	Antibodies Detection Rapid Test Kit			
<b>Product Specification for BIOGENIX (BSURE)--BIOGENIX INC. PRIVATE LIMITED HIV1 &amp; HIV2 Antibodies Detection Rapid Test Kit</b>				
<b>विवरण Specification</b>	<b>उप-विवरण 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/2020/000068	
	Manufacturing unit certification		ISO:13485 (Latest)	
	Additional voluntary certification available		ISO:9001 , QMS	
	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
इंपीबीजी विवरण   ePBG Detail		
NA		
<b>नियम और शर्तें  Terms and Conditions</b>		
<p><b>1. Special terms and conditions- Version:2 effective from 19-05-2023</b></p> <p>1.1</p> <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> <p><b>2. General Terms and Conditions-</b></p> <p>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.</p> <p>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</p>		
<p>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।</p> <p>Note: This is system generated file. No signature is required. Print out of this document is not valid for payment/ transaction purpose.</p>		