

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 Designation of Administrative Approval:	cmo lalitpur	भुगतान का तरीका Payment Mode:	Offline			
वित्तीय अनुमोदन का पदनाम Designation of Financial Approval :	cmo lalitpur	पद Designation :	ACMO			
		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 CHAURAHA,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 चतुर्थांश 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				
Product Specification for BIOGENIX (BSURE)--BIOGENIX INC. PRIVATE LIMITED HIV1 & HIV2 Antibodies Detection Rapid Test Kit						
विनिर्देश Specification	उप-विनिर्देश Sub-Spec				मूल्य Value	
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 specificy 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
ईपीबीजी विवरण ePBG Detail		
NA		
नियम और शर्तें Terms and Conditions		
<p>1. Special terms and conditions- Version:2 effective from 19-05-2023</p> <p>1.1</p> <ol style="list-style-type: none"> 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 self-declaration of valid Drug License, product certification, test reports etc. However, buyers must check and validate the details at their end for all applicable licenses and certifications e.g., validity and authenticity/genuineness of drug license, product certification, manufacturer certification/licenses, test reports etc. 3. 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 & regulatory liability in respect of the offered/supplied product shall be totally of both manufacturer and reseller/distributor. 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. <p>2. General Terms and Conditions-</p> <p>2.1 This contract is governed by the General Terms and Conditions, 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>		