

अनुबंध | Contract



अनुबंध क्रमांक | Contract No: GEMC-511687795133930

अनुबंध तिथि | Generated Date : 22-Mar-2024

संगठन विवरण Organisation Details	खरीदार विवरण Buyer Details
प्ररूप Type : State Government मंत्रालय Ministry : - विभाग Department : Health Medical Education and Family Welfare Department Jharkhand संगठन का नाम Organisation Name : N/A कार्यालय क्षेत्र Office Zone: CIVIL SURGEON OFFICE GODDA	पद Designation : LAURENTUS TIRKEY संपर्क नंबर Contact No. : 06422-000000- ईमेल आईडी Email ID : buycon25.hmefwdj.jh@gembuyer.in जीएसटीआईएन GSTIN : - पता Address : CIVIL SURGEON OFFICE, GODDA, PO GODDA ,PS GODDA, DISTRICT GODDA ,JHARKHAND, GODDA, JHARKHAND-814133, 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: CIVIL SURGEON GODDA वित्तीय अनुमोदन का पदनाम Designation of Financial Approval : CIVIL SURGEON GODDA	Role: BUYER भुगतान का तरीका Payment Mode: Offline पद Designation : LAURENTUS TIRKEY ईमेल आईडी Email ID : buycon25.hmefwdj.jh@gembuyer.in जीएसटीआईएन GSTIN : - पता Address: CIVIL SURGEON OFFICE, GODDA, PO GODDA ,PS GODDA, DISTRICT GODDA ,JHARKHAND, Godda, JHARKHAND-814133, India

विक्रेता विवरण Seller Details
जेम विक्रेता आईडी GeM Seller ID : HYEF210003147141 कंपनी का नाम Company Name : M/S. CHANDRA IMPEX संपर्क नंबर Contact No. : 07250555353 ईमेल आईडी Email ID : goddachandrainpex@gmail.com पता Address : 319,CHANDRA IMPEX,GODDA,MISSION CHOWK, Godda, JHARKHAND-814133, - एमएसएमई पंजीकरण संख्या MSME Registration number : udyam-jh-09-0001281 एमएसई सामाजिक श्रेणी MSE Social Category : OBC एमएसई लिंग श्रेणी MSE Gender : Male जीएसटीआईएन GSTIN: 20AVCPK4419Q1ZI

*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 : PAREEKSHAK Dual Rapid Test Kit for HIV and Syphilis ब्रांड Brand : PAREEKSHAK 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 : Dual Rapid Test Kit for HIV and Syphilis (Q2) मॉडल Model: H12SWB-50 एचएसएन कोड HSN Code: HSN not specified by seller	7,800	Test	35	NA	273,000
कुल ऑर्डर मूल्य Total Order Value (in INR)						273,000

परोक्षी विवरण Consignee Detail						
क्र.सं. S.No	परोक्षी 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	संपर्क Contact : 06422-000000- जीएसटीआईएन GSTIN : - पता Address : CIVIL SURGEON OFFICE, GODDA, PO GODDA ,PS GODDA, DISTRICT GODDA ,JHARKHAND, GODDA, JHARKHAND-814133, India	PAREEKSHAK Dual Rapid Test Kit for HIV and Syphilis	-	7,800	22-Mar-2024	06-Apr-2024
Product Specification for PAREEKSHAK Dual Rapid Test Kit for HIV and Syphilis						
विनिर्देश Specification	उप-विनिर्देश Sub-Spec	मूल्य Value				
GENERAL FEATURES	Product Description	Dual Rapid Test Kit for HIV and Syphilis				
	Clinical Purpose	For in-vitro detection of HIV and Syphilis infection				
PRODUCT INFORMATION	Assay type	Synthetic and/or recombinant antigens coated on solid phase a) Multiple Treponema pallidum antigens, and b) Multiple Antigens of HIV1 (including gp41) and HIV2 (including gp36)				
	Detects	Individually detects antibodies to all subtypes of HIV1 & HIV2 and Treponema pallidum by the rapid test, in human whole blood/ & serum/ & plasma				
	Detection Type	Qualitative				
	Testing Principle	Lateral Flow (Immunochromatography)				
	Species Reactivity	Human				
	Type of Sample	Whole Blood,Serum,Plasma				
	Time to Result	≤ 30 minutes				
	Ability to evaluate negative or positive test result	Yes				
	Assay Sensitivity for HIV	≥ 100%				
	Assay Specificity for HIV	≥ 98%				
	Assay Sensitivity for Syphilis	≥ 85%				
	Assay Specificity for Syphilis	≥ 93%				
	Declared sensitivity and specificity shall be claimed by the manufacturer in the kit literature	Yes				
	Contains an internal control dot/band (in-built control) for the confirmation that the test has been performed correctly	Yes				
	The control dot/band (in-built control) 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	No				
KIT CONTENTS	Kit Contents	Test Device/Card/Cassette with Desiccant, Sample Dropper, Assay Buffer (if any)				
	The kit shall be supplied with sufficient amount of capillaries to deliver the required amount of specimen specified in the kit literature	Yes				
	Positive and negative controls provided with each pack of kit	No				
	Quantity of positive and negative controls provided	NA				
	Documents detailing the principal, components, details of antigen used/coated for detection of HIV 1&2 as well as Treponema pallidum antibodies, biosafety compliance, validity criteria, interpretation of results, performance characteristic provided	Yes				
	Manufacturing and expiry dates should be mentioned for each kit	Yes				
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/2020/000013				
	Manufacturing unit certification	ISO:13485 (Latest)				
	Additional voluntary certification available	ISO 9001:2015				
	Availability of Test Report for each supplied batch/product as per Medical Device Rules	Yes				

	(MDR) 2017 as amended till date	
	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

1. Special terms and conditions- Version:2 effective from 19-05-2023

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.
- 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.
- 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 drug license held by them.
- 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.
- 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.

2. General Terms and Conditions-

- This contract is governed by the [General Terms and Conditions](#), conditions stipulated to this Product/Service as provided in the Marketplace.
- 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.