

## अनुबंध | Contract



अनुबंध क्रमांक | Contract No: GEMC-511687722940996

अनुबंध तिथि | Generated Date : 14-Jun-2024

संगठन विवरण   Organisation Details	खरीदार विवरण   Buyer Details
प्ररूप   Type : State Government मंत्रालय   Ministry : - विभाग   Department : Medical Health and Family Welfare Department Uttar Pradesh संगठन का नाम   Organisation Name : N/A कार्यालय क्षेत्र   Office Zone: Agra	पद   Designation : MEDICAL SUPERINTENDENT संपर्क नंबर   Contact No. : 0562-2446099- ईमेल आईडी   Email ID : nodaldmh.ag-up@gov.in जीएसटीआईएन   GSTIN : 09AAAGD0871P1DC पता   Address : District Hospital Agra, MG Road darsh nagar chhipitola rakabganj Agra, AGRA, UTTAR PRADESH-282001, 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: SIC DH Agra वित्तीय अनुमोदन का पदनाम   Designation of Financial Approval : SIC DH Agra	Role: BUYER भुगतान का तरीका   Payment Mode: Offline पद   Designation : MEDICAL SUPERINTENDENT ईमेल आईडी   Email ID : nodaldmh.ag-up@gov.in जीएसटीआईएन   GSTIN : 09AAAGD0871P1DC पता   Address: District Hospital Agra, MG Road darsh nagar chhipitola rakabganj Agra, AGRA, UTTAR PRADESH-282001, India

विक्रेता विवरण   Seller Details
जेम विक्रेता आईडी   GeM Seller ID : 49D7180000115801 कंपनी का नाम   Company Name : K S PHARMA संपर्क नंबर   Contact No. : 09927350818 ईमेल आईडी   Email ID : shailendrasingh001@gmail.com पता   Address : 42/99, BILLOCHPURA, MADIYA KATRA, Agra, UTTAR PRADESH-282002, - एमएसएमई पंजीकरण संख्या   MSME Registration number : UDYAM-UP-01-0017108 एमएसई सामाजिक श्रेणी   MSE Social Category : General एमएसई लिंग श्रेणी   MSE Gender : Male जीएसटीआईएन   GSTIN: 09BMGPS1929H1ZP (R)

\*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 : SIDAK LIFE CARE HCV Total Antibodies Detection Rapid Test Kit ब्रांड   Brand : SIDAK LIFE CARE 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 : HCV Rapid Test Kits (Q2) मॉडल   Model: SIDAK LIFE CARE एचएसएन कोड   HSN Code: HSN not specified by seller	500	Test	49.95	NA	24,975
कुल ऑर्डर मूल्य   Total Order Value (in INR)						24,975

परोक्ष विवरण   Consignee Detail						
क्र.सं.   S.No	परोक्ष   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 : BUYER-CONSIGNEE ईमेल आईडी   Email ID : nodaldmh.ag-up@gov.in					

1	संपर्क   Contact : 0562-2446099- जीएसटीआईएन   GSTIN : 09AAAGD0871P1DC पता   Address : District Hospital Agra, MG Road darsh nagar chhipitola rakabganj Agra, AGRA, UTTAR PRADESH-282001, India	SIDAK LIFE CARE HCV Total Antibodies Detection Rapid Test Kit	-	500	14-Jun-2024	29-Jun-2024
<b>Product Specification for SIDAK LIFE CARE HCV Total Antibodies Detection Rapid Test Kit</b>						
<b>विनिर्देश   Specification</b>	<b>उप-विनिर्देश   Sub-Spec</b>	<b>मूल्य   Value</b>				
GENERAL FEATURES	Product Description	HCV Rapid Test Kit				
	Clinical Purpose	To provide diagnosis of Hepatitis C Virus infection				
PRODUCT INFORMATION	Type of Kit	HCV Total Antibodies Detection Rapid Test Kit				
	Generation	3rd Generation				
	Solid phase/particle coated with recombinant and / or synthetic peptide antigens for Core, NS3, NS 4, and NS5	Yes				
	Detection Type	Qualitative				
	Testing Principle	Lateral Flow Immunochromatographic Assay				
	Species Reactivity	Human				
	Type of Sample	Serum, Plasma				
	Time to Result	≤ 30 minutes				
	Ability to evaluate negative or positive test result	Yes				
	Sensitivity	≥95%				
	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	No					
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				
	Adequate document detailing principle, components, methodologies, validity criteria, interpretation of results, performance characteristics, bio-safety, limitations of assay, storage condition, mfg & exp date and method of disposal provided with kit	Yes				
	Positive and negative controls provided with each pack of kit	No				
	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/2021/000009				
	Manufacturing unit certification	ISO:13485 (Latest)				
	Additional voluntary certification available	UQ-2021020570				
	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				
	Agree to provide advance sample of the product for buyer's approval before commencement of supply in case of					

ADVANCE SAMPLE	bidding	No
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		
<p>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.</p> <p>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.</p> <p>3. In case of authorized resellers/distributors, it will be the legal &amp; 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.</p> <p>4. The prices 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.</p> <p>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-		
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		
<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>		