

## अनुबंध | Contract



अनुबंध क्रमांक | Contract No: GEMC-511687712341160

अनुबंध तिथि | Generated Date : 19-Jun-2024

संगठन विवरण   Organisation Details		खरीदार विवरण   Buyer Details				
प्ररूप   Type :	State Government	पद   Designation :	ACMO RCH			
मंत्रालय   Ministry :	-	संपर्क नंबर   Contact No. :	-			
विभाग   Department :	Medical Health and Family Welfare Department Uttar Pradesh	ईमेल आईडी   Email ID :	buycon352.mhfwdup.up@gembuyer.in			
संगठन का नाम   Organisation Name :	N/A	जीएसटीआईएन   GSTIN :	-			
कार्यालय क्षेत्र   Office Zone:	Chief Medical Officer Fatehpur	पता   Address :	Office of Chief Medical Officer, Turab Ali Ka Purwa, Fatehpur - 212601, FATEHPUR, UTTAR PRADESH-212601, 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	भुगतान का तरीका   Payment Mode:	Offline			
वित्तीय अनुमोदन का पदनाम   Designation of Financial Approval :	cmo	पद   Designation :	ACMO RCH			
		ईमेल आईडी   Email ID :	buycon352.mhfwdup.up@gembuyer.in			
		जीएसटीआईएन   GSTIN :	-			
		पता   Address:	Office of Chief Medical Officer, Turab Ali Ka Purwa, Fatehpur - 212601, Fatehpur, UTTAR PRADESH-212601, India			
विक्रेता विवरण   Seller Details						
जेम विक्रेता आईडी   GeM Seller ID :	A616190000822480					
कंपनी का नाम   Company Name :	M/S NIDHI TRADING COMPANY					
संपर्क नंबर   Contact No. :	09415041818					
ईमेल आईडी   Email ID :	nidhitradingco.kanpur@gmail.com					
पता   Address :	117/290, N-BLOCK,KAKADEV, KANPUR, UTTAR PRADESH-208011, -					
एमएसएमई पंजीकरण संख्या   MSME Registration number :	UDYAM-UP-43-0008817					
एमएसई सामाजिक श्रेणी   MSE Social Category :	General					
एमएसई लिंग श्रेणी   MSE Gender :	Male					
जीएसटीआईएन   GSTIN:	09ABSPB8812A1ZS (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 : AVECON--AVECON HEALTHCARE PVT LTD Salmonella Typhi IgM Antibody Detection Rapid Test Kit ब्रांड   Brand : AVECON--AVECON HEALTHCARE PVT LTD 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 : Typhoid Rapid Test Kits (Q2) मॉडल   Model: MAXLINE TYPHOID ANTIBODY IgM-IgG (50 TEST) एचएसएन कोड   HSN Code: HSN not specified by seller	2,400	Test	55	NA	132,000
कुल ऑर्डर मूल्य   Total Order Value (in INR)						132,000
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
	पद   Designation : Chief Pharmacist ईमेल आईडी   Email ID : arimardan.05787@gov.in	AVECON--AVECON				

1	संपर्क   Contact : - जीएसटीआईएन   GSTIN : - पता   Address : Office of Chief Medical Officer, Turab Ali Ka Purwa, Fatehpur - 212601, FATEHPUR, UTTAR PRADESH-212601, India	HEALTHCARE PVT LTD Salmonella Typhi IgM Antibody Detection Rapid Test Kit	-	2,400	19-Jun-2024	04-Jul-2024
Product Specification for AVECON--AVECON HEALTHCARE PVT LTD Salmonella Typhi IgM Antibody Detection Rapid Test Kit						
विनिर्देश   Specification	उप-विनिर्देश   Sub-Spec			मूल्य   Value		
GENERAL FEATURES	Product Description			Typhoid Rapid Test Kit		
	Clinical Purpose			To detect the presence of antibodies associated with Salmonella typhi in human blood samples		
PRODUCT INFORMATION	Type of Kit			Salmonella Typhi IgM Antibody Detection Rapid Test Kit		
	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			≥98%		
	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		
	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			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 Rule (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/2019/000041		
	Manufacturing unit certification			ISO:13485 (Latest)		
	Additional voluntary certification available			UQ-14088		
	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		
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 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.</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.		