

## अनुबंध | Contract



अनुबंध क्रमांक | Contract No: GEMC-511687723950191

अनुबंध तिथि | Generated Date : 30-Mar-2024

<b>संगठन विवरण   Organisation Details</b>		<b>खरीदार विवरण   Buyer Details</b>				
प्ररूप   Type :	State Government	पद   Designation :	MO			
मंत्रालय   Ministry :	-	संपर्क नंबर   Contact No. :	02772-275316-			
विभाग   Department :	Panchayats and Rural Housing Department Gujarat	ईमेल आईडी   Email ID :	buycon354.prhdg.gj@gembuyer.in			
संगठन का नाम   Organisation Name :	N/A	जीएसटीआईएन   GSTIN :	-			
कार्यालय क्षेत्र   Office Zone:	Arvalli District Panchayat	पता   Address :	Chorimala CHC, SABARKANTHA, GUJARAT-383246, India			
<b>वित्तीय स्वीकृति विवरण   Financial Approval Detail</b>		<b>भुगतान प्राधिकरण विवरण   Paying Authority Details</b>				
आईएफडी सहमति   IFD Concurrence :	No	Role:	PAO			
प्रशासनिक अनुमोदन का पदनाम   Designation of Administrative Approval:	MO	भुगतान का तरीका   Payment Mode:	Internet Banking			
वित्तीय अनुमोदन का पदनाम   Designation of Financial Approval :	MO	पद   Designation :	AO Health			
		ईमेल आईडी   Email ID :	ao-mod-health-arv@gujarat.gov.in			
		जीएसटीआईएन   GSTIN :	-			
		पता   Address:	District Panchayat Arvalli, Sabarkantha, GUJARAT-383315, India			
<b>विक्रेता विवरण   Seller Details</b>						
जेम विक्रेता आईडी   GeM Seller ID :	6JTP220005998295					
कंपनी का नाम   Company Name :	DARSHAN DJAGNOSTIC					
संपर्क नंबर   Contact No. :	08320052739					
ईमेल आईडी   Email ID :	mehtacool_1970@yahoo.co.in					
पता   Address :	B2,BALAJI COMPLEX,MALPUR ROAD,OPP. TALUKA PANCHAYAT, Sabarkantha, GUJARAT-383315, -					
एमएसएमई पंजीकरण संख्या   MSME Registration number :	-					
जीएसटीआईएन   GSTIN:	24AOKPM2791J1ZL					
*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						
<b>उत्पाद विवरण   Product Details</b>						
#	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 Syphilis Antibody 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 : Syphilis Rapid Test Kits (Q2) मॉडल   Model: MAXLINE SYPHILIS CARDS (WB) (50TEST) एचएसएन कोड   HSN Code: HSN not specified by seller	50	Test	14	NA	700
कुल ऑर्डर मूल्य   Total Order Value (in INR)						700
<b>परोक्ष विवरण   Consignee Detail</b>						
क्र.सं.   S.No	परोक्ष विवरण   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 : MO ईमेल आईडी   Email ID : buycon354.prhdg.gj@gembuyer.in संपर्क   Contact : 02772-275316- जीएसटीआईएन   GSTIN : - पता   Address : Chorimala CHC, SABARKANTHA, GUJARAT-383246, India	AVECON--AVECON HEALTHCARE PVT LTD Syphilis Antibody Rapid Test Kit	-	50	30-Mar-2024	14-Apr-2024

**Product Specification for AVECON--AVECON HEALTHCARE PVT LTD Syphilis Antibody Rapid Test Kit**

विनिर्देश   Specification	उप-विनिर्देश   Sub-Spec	मूल्य   Value
GENERAL FEATURES	Product Description	Syphilis Rapid Test Kit
	Clinical Purpose	To provide a quick and accurate diagnosis of syphilis infection in all stages of infection by detecting antibodies to Treponema Pallidum
PRODUCT INFORMATION	Type of Kit	Syphilis Antibody Rapid Test Kit
	Detects	Total Anti-Treponema Pallidum Antibody (IgG,IgM & IgA)
	Detection Type	Qualitative
	The assay shall have solid phase coated with synthetic or recombinant type of Treponema Pallidum antigens	Yes
	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	≥99%
	Specificity	≥98%
	Declared sensitivity and specificity shall be claimed by the manufacturer in the kit literature	Yes
	The assay calibrated to WHO reference serum and the same shall be supported by statements in kit insert and certificate from the manufacturer	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	NA
	Each test card/strip supplied with sterile auto retractable disposable lancet and disposable alcoholic swabs	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 with 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 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
	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
<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 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.</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		
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.		