

## Contract | अनुबंध



Contract No | अनुबंध क्रमांक: GEMC-511687732949678

Generated Date | अनुबंध तिथि: 29-Dec-2023

Organisation Details   संगठन विवरण		Buyer Details   खरीदार विवरण	
Type   प्रकृति :	State Government	Designation   पद :	ACCOUNT MANAGER
Ministry   मंत्रालय :	-	Contact No.   संपर्क नंबर :	01995-261410-
Department   विभाग :	Health and Family Welfare Department Jammu and Kashmir	Email ID   ईमेल आईडी :	buyer33.hfwdjk.jk@gembuyer.in
Organisation Name   संगठन का नाम :	N/A	GSTIN   जीएसटीआईएन :	-
Office Zone   कार्यालय क्षेत्र:	kishtwar	Address   पता :	District Hospital Kishtwar, DODA, JAMMU & KASHMIR-182204, India

Financial Approval Detail   वित्तीय स्वीकृति विवरण		Paying Authority Details   भुगतान प्राधिकरण विवरण	
IFD Concurrence   आईएफडी सहमति:	No	Role:	BUYER
Designation of Administrative Approval   प्रशासनिक अनुमोदन का पदनाम:	MEDICAL SUPERINTENDENT DISTRICT HOSPITAL K ISHTWAR	Payment Mode   भुगतान का तरीका:	Offline
Designation of Financial Approval   वित्तीय अनुमोदन का पदनाम :	MEDICAL SUPERINTENDENT DISTRICT HOSPITAL K ISHTWAR	Designation   पद :	ACCOUNT MANAGER
		Email ID   ईमेल आईडी :	buyer33.hfwdjk.jk@gembuyer.in
		GSTIN   जीएसटीआईएन :	-
		Address   पता:	District Hospital Kishtwar, Kishtwar, JAMMU & KASHMIR-182204, India

Seller Details   विक्रेता विवरण	
GeM Seller ID   जेम विक्रेता आईडी :	B096190000823475
Company Name   कंपनी का नाम :	JAMMU ENTERPRISES
Contact No.   संपर्क नंबर :	09419144530
Email ID   ईमेल आईडी :	jammuenterprises82@gmail.com
Address   पता :	,90,MAHESHPURA JAMMU, JAMMU, JAMMU AND KASHMIR-180001, -
MSME Registration number   एमएसएमई पंजीकरण संख्या :	-
GSTIN   जीएसटीआईएन:	01AAKFJ9839L1ZV

\*GST / Tax invoice to be raised in the name of | जिसके नाम के पक्ष में GST/TAX इनवॉइस पेश किया जाएगा - Buyer

Delivery Instructions   वितरण निर्देश : NA	
Product Details   उत्पाद विवरण	

#	Item Description   आइटम विवरण	Ordered Quantity   आइटम विवरण	Unit   इकाई	Unit Price (INR)   इकाई मूल्य (INR)	Tax Bifurcation (INR)   कर विभाजन (INR)	Price (Inclusive of all Duties and Taxes in INR)   मूल्य (INR में सभी शुल्क और कर सहित)
1	Product Name   उत्पाद का नाम : ErbaQik Syphilis Antibody Rapid Test Kit Brand   ब्रांड : ErbaQik Brand Type   ब्रांड प्रकार : Registered Brand Catalogue Status   कैटलॉग की स्थिति: OEM verified catalogue Selling As   कैसे बेचा जा रहा है : OEM verified Reseller Category Name & Quadrant   श्रेणी का नाम और चतुर्धारा : Syphilis Rapid Test Kits (Q2) Model   मॉडल: ErbaQik Syphilis Test Card (25 Test ) HSN Code   एचएसएन कोड: HSN not specified by seller	200	Test	17.88	NA	3,576
Total Order Value   कुल ऑर्डर मूल्य (in INR)						3,576

Consignee Detail   प्रेषिती विवरण						
S.No   क्र.सं.	Consignee   प्रेषिती	Item   वस्तु	Lot No.   लॉट नंबर	Quantity   मात्रा	Delivery Start After   दिनांक के बाद डिलीवरी शुरू करना है	Delivery To Be Completed By   वितरण पूरा कब तक करना है
1	Designation   पद : Primary User Email ID   ईमेल आईडी : ms-kishtwar@jk.gov.in Contact   संपर्क : 01995-261410- GSTIN   जीएसटीआईएन : - Address   पता : District Hospital Kishtwar, DODA, JAMMU & KASHMIR-182204, India	ErbaQik Syphilis Antibody Rapid Test Kit	-	200	29-Dec-2023	13-Jan-2024

Product Specification for ErbaQik Syphilis Antibody Rapid Test Kit		
Specification   विवरण	Sub-Spec   उप-विवरण	Value   मूल्य
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
KIT CONTENTS	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	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
PACKAGING	Quantity of positive and negative controls provided	NA
	Each test card/strip supplied with sterile auto retractable disposable lancet and disposable alcoholic swabs	No
	Pack size of kit	25 Tests
CERTIFICATIONS & REPORTS	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
	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/2020/000014
	Manufacturing unit certification	ISO:13485 (Latest)
	Additional voluntary certification available	ISO 9001
SHELF LIFE	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

## ePBG Detail | ईपीबीजी विवरण

NA

### Terms and Conditions | नियम और शर्तें

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

1.1

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.
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 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.
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.

#### 2. General Terms and Conditions-

2.1 This contract is governed by the [General Terms and Conditions](#), 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

Note: This is system generated file. No signature is required. Print out of this document is not valid for payment/ transaction purpose.

नोट: यह सिस्टम जनरेटेड फाइल है। कोई हस्ताक्षर की आवश्यकता नहीं है। इस दस्तावेज़ का प्रिटे आउट भुगतान/लेनदेन उद्देश्य के लिए मान्य नहीं है।