

अनुबंध | Contract



अनुबंध क्रमांक | Contract No: GEMC-511687782154583

अनुबंध तिथि | Generated Date : 28-Mar-2024

संगठन विवरण | Organisation Details

प्ररूप | Type : State Government
मंत्रालय | Ministry : -
विभाग | Department : Health and Family Welfare Department Delhi
संगठन का नाम | Organisation Name : N/A
कार्यालय क्षेत्र | Office Zone: Lok Nayak Hospital

खरीदार विवरण | Buyer Details

पद | Designation : link to procurement officer
संपर्क नंबर | Contact No. : 011-23501229-
ईमेल आईडी | Email ID : dr.ankaj@delhi.gov.in
जीएसटीआईएन | GSTIN : 07DELL02991C1DE
पता | Address : Lok Nayak Hospital, Jawahar Lal Nehru Marg, New Delhi-110002,
CENTRAL DELHI, DELHI-110002, India

वित्तीय स्वीकृति विवरण | Financial Approval Detail

आईएफडी
सहमति | IFD No
Concurrence :
प्रशासनिक
अनुमोदन का
पदनाम |
Designation MD, LNH
of
Administrative
Approval:
वित्तीय अनुमोदन
का पदनाम | Medical Director Head Major Head 2210 Medical & Public Health, 01 Ur
Designation ban Health Services 110 Hospital & Dispensaries 86- Lok Nayak Hospital
of Financial al 2210-01-110-86-0021 Material & Supplies under Demand No 07 for t
Approval: he financial Year 2023-24

भुगतान प्राधिकरण विवरण | Paying Authority Details

Role: PAO
भुगतान का तरीका |
Payment Mode: Offline
पद | Designation : DDO
ईमेल आईडी | Email ID : varun.rahal45@delhi.gov.in
जीएसटीआईएन | GSTIN : -
पता | Address: Lok Nayak Hospital, Jawahar Lal Nehru Marg, New Delhi-110002 9873617042,
CENTRAL DELHI, DELHI-110002, India

विक्रेता विवरण | Seller Details

जेम विक्रेता आईडी | GeM Seller ID : 15B4180000117531
कंपनी का नाम | Company Name : Q-LINE BIOTECH PRIVATE LIMITED
संपर्क नंबर | Contact No. : 09415022090
ईमेल आईडी | Email ID : poctlko@pocservices.com
पता | Address : 298-281,SERVICES,298 281 Transport Nagar Kanpur Road Adjacent Transport Nagar Metro Station Lucknow 226023,84000,
Lucknow, UTTAR PRADESH-226023, India
एमएसएमई पंजीकरण संख्या | MSME Registration number : -
जीएसटीआईएन | GSTIN: 09AAFCP9672B1Z6

*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 : Q-line HBsAg Rapid Test Kit ब्रांड Brand : Q-line ब्रांड प्रकार Brand Type : Registered Brand कैटलॉग की स्थिति Catalogue Status: OEM verified catalogue कैसे बेचा जा रहा है Selling As : OEM श्रेणी का नाम और चतुर्थांश Category Name & Quadrant : HBsAg Rapid Test Kits (Q2) मॉडल Model: HBS-025PS एचएसएन कोड HSN Code: HSN not specified by seller	1,500	Test	8.28	NA	12,420
कुल ऑर्डर मूल्य Total Order Value (in INR)						12,420

परोक्षिती विवरण | Consignee Detail

क्र.सं. S.No	परोक्षिती Consignee	वस्तु Item	लॉट नंबर Lot No.	मात्रा Quantity	दिनांक के बाद डिलीवरी शुरू करना है Delivery Start	वितरण पूरा कब तक करना है Delivery To Be Completed
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1	<p>पद Designation : MOI/C- Equipment Store ईमेल आईडी Email ID : sujoy.neogi@gov.in संपर्क Contact : 011-23232400-1312 जीएसटीआईएन GSTIN : - पता Address : Lok Nayak Hospital, Jawahar Lal Nehru Marg, New Delhi-110002, CENTRAL DELHI, DELHI-110002, India</p>	Q-line HBsAg Rapid Test Kit	-	1,500	28-Mar-2024	12-Apr-2024
Product Specification for Q-line HBsAg Rapid Test Kit						
विनिर्देश Specification	उप-विनिर्देश Sub-Spec				मूल्य Value	
GENERAL FEATURES	Product Description				HBsAg Rapid Test Kit	
	Clinical Purpose				To provide diagnosis of Hepatitis B Virus infection	
PRODUCT INFORMATION	Detects				Hepatitis B Surface Antigen (HBsAg)	
	Should be solid phase/ particle coated with monoclonal antibodies to HBsAg				Yes	
	Test Should be able to detect all 11 subtype and Variants of HBsAg				Yes	
	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	
	Assay Sensitivity (%)				100%	
	Assay 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				25 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				UP3220B003905	
	Manufacturing unit certification				ISO:13485 (Latest)	
	Additional voluntary certification available				CE Certificate	
	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	3/4th 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:1 effective from 04-05-2023

- 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-

- 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

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।

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