

अनुबंध | Contract



अनुबंध क्रमांक | Contract No: GEMC-511687792215891

अनुबंध तिथि | Generated Date : 29-May-2024

संगठन विवरण | Organisation Details

प्ररूप | Type : State Government
मंत्रालय | Ministry : -
विभाग | Department : Health and Family Welfare Department Delhi
संगठन का नाम | Organisation Name : Directorate General of Health Services
कार्यालय क्षेत्र | Office Zone : Guru Nanak Eye Centre

खरीदार विवरण | Buyer Details

पद | Designation : purchase officer
संपर्क नंबर | Contact No. : 011-23234622-285
ईमेल आईडी | Email ID : dr.paromitadutta@delhi.gov.in
जीएसटीआईएन | GSTIN : 07DELG00429C1D4
पता | Address : Maharaja Ranjit Singh Marg,
CENTRAL DELHI, DELHI-110002, India

वित्तीय स्वीकृति विवरण | Financial Approval Detail

आईएफडी सहमति | IFD Concurrence : No
प्रशासनिक अनुमोदन का पदनाम | Designation of Administrative Approval : Director, GNEC
वित्तीय अनुमोदन का पदनाम | Designation of Financial Approval : Director, GNEC

भुगतान प्राधिकरण विवरण | Paying Authority Details

Role: BUYER
भुगतान का तरीका | Payment Mode: Offline
पद | Designation : purchase officer
ईमेल आईडी | Email ID : dr.paromitadutta@delhi.gov.in
जीएसटीआईएन | GSTIN : 07DELG00429C1D4
पता | Address : Maharaja Ranjit Singh Marg,
CENTRAL DELHI, DELHI-110002, India

विक्रेता विवरण | Seller Details

जेम विक्रेता आईडी | GeM Seller ID : RVGM230008150848
कंपनी का नाम | Company Name : M/s Sunshine Pharma
संपर्क नंबर | Contact No. : 09873843669
ईमेल आईडी | Email ID : sunshinepharma3012@gmail.com
पता | Address : Block -A,A-282, Shastri Nagar, New Delhi, Central Delhi, Delhi, 110052, New Delhi, Central Delhi, Delhi, 110052,
North West delhi, DELHI-110052, -
एमएसएमई पंजीकरण संख्या | MSME Registration number : -
जीएसटीआईएन | GSTIN : 07APNPG2931A2ZA (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 : PK SIMPLE HCV Total Antibodies Detection Rapid Test Kit ब्रांड Brand : PK SIMPLE 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: PKRK013 एचएसएन कोड HSN Code: HSN not specified by seller | 100 | Test | 13 | NA | 1,300 |
| कुल ऑर्डर मूल्य Total Order Value (in INR) | | | | | | 1,300 |

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 |
|----------------|--|---|--------------------|-------------------|---|--|
| 1 | पद Designation : MO Incharge ईमेल आईडी Email ID : m.wadhvani08@delhi.gov.in संपर्क Contact : 011-23239115-262 जीएसटीआईएन GSTIN : - पता Address : Maharaja Ranjit Singh Marg, CENTRAL DELHI, DELHI-110002, India | PK SIMPLE HCV Total Antibodies Detection Rapid Test Kit | - | 100 | 29-May-2024 | 13-Jun-2024 |

Product Specification for PK SIMPLE HCV Total Antibodies Detection Rapid Test Kit

| विनिर्देश Specification | उप-विनिर्देश Sub-Spec | मूल्य Value |
|---------------------------|---|---|
| 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 | 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 |
| | 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 | MFG/IVD/2021/000068 |
| | Manufacturing unit certification | ISO:13485 (Latest) |
| | Additional voluntary certification available | NA |
| | 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 |

NA

नियम और शर्तें | Terms and Conditions

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

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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