# अनुबंध|Contract

**अनुबंध क्रमांक|Contract No:** GEMC-511687765265409
**अनुबंध तिथि|Generated Date :** 14-Mar-2024

## संगठन विवरण|Organisation Details

| | |
|---|---|
| प्ररूप|Type : | State Government |
| मंत्रालय|Ministry : | - |
| विभाग|Department : | Medical Health and Family Welfare Department Uttar Pradesh |
| संगठन का नाम|Organisation Name : | N/A |
| कार्यालय क्षेत्र|Office Zone: | BRD Hospital MahanagarLucknow |

## खरीदार विवरण|Buyer Details

| | |
|---|---|
| पद|Designation : | Medical Superintendent |
| संपर्क नंबर|Contact No. : | 0522-9305012-069 |
| ईमेल आईडी|Email ID : | cms.brdh.lu-up@gov.in |
| जीएसटीआईएन|GSTIN : | 09LKNC00830E1DK |
| पता|Address : | BRD Hospital, Mahanagar, Lucknow-226010, LUCKNOW, UTTAR PRADESH-226010, India |

## वित्तीय स्वीकृति विवरण|Financial Approval Detail

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|---|---|
| आईएफडी सहमति|IFD Concurrence : | No |
| प्रशासनिक अनुमोदन का पदनाम\| Designation of Administrative Approval: | CMS |
| वित्तीय अनुमोदन का पदनाम\| Designation of Financial Approval : | CMS |

## भुगतान प्राधिकरण विवरण|Paying Authority Details

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|---|---|
| Role: | BUYER |
| भुगतान का तरीका\| Payment Mode: | Offline |
| पद|Designation : | Medical Superintendent |
| ईमेल आईडी|Email ID : | cms.brdh.lu-up@gov.in |
| जीएसटीआईएन|GSTIN : | 09LKNC00830E1DK |
| पता|Address: | BRD Hospital, Mahanagar, Lucknow-226010, Lucknow, UTTAR PRADESH-226010, India |

## विक्रेता विवरण|Seller Details

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|---|---|
| जेम विक्रेता आईडी|GeM Seller ID : | ISJM210003835149 |
| कंपनी का नाम|Company Name : | NEELIMA ENTERPRISES |
| संपर्क नंबर|Contact No. : | 09208583808 |
| ईमेल आईडी|Email ID : | neelima.enterprises009@gmail.com |
| पता\| Address : | S/O JAGANNATH PRASAD, SIDHAULI,NAROTTAM NAGAR,SIDHAULI,SIDHAULI, Sitapur, UTTAR PRADESH-261303, - |
| एमएसएमई पंजीकरण संख्या|MSME Registration number : | - |
| जीएसटीआईएन|GSTIN: | 09BICPK1757M1ZW |

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

वितरण निर्देश | Delivery Instructions : Good Quality and Lower Price

## 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 :** Merilisa Dengue IgM + IgG Antibody Detection Rapid Test Kit<br>**ब्रांड\|Brand :** Merilisa<br>**ब्रांड प्रकार\|Brand Type :** Registered Brand<br>**कैटलॉग की स्थिति\|Catalogue Status:** OEM verified catalogue<br>**कैसे बेचा जा रहा है\|Selling As :** OEM verified Reseller<br>**श्रेणी का नाम और चतुर्थांश\|Category Name & Quadrant :** Dengue Rapid Test Kits (Q2)<br>**मॉडल\|Model:** MeriScreen Dengue IgG IgM<br>**एचएसएन कोड\|HSN Code:** HSN not specified by seller | 1,000 | Test | 240 | NA | 240,000 |
| कुल ऑर्डर मूल्य \|Total Order Value (in INR) | | | | | | **240,000** |

## 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 |
|---|---|---|---|---|---|---|
| 1 | पद\| Designation : Medical Superintendent<br>ईमेल आईडी\|Email ID : cms.brdh.lu-up@gov.in<br>संपर्क\|Contact : 0522-9305012-069<br>जीएसटीआईएन\|GSTIN : 09LKNC00830E1DK<br>पता\|Address : BRD Hospital, Mahanagar, Lucknow-226010, LUCKNOW, UTTAR PRADESH-226010, India | Merilisa Dengue IgM + IgG Antibody Detection Rapid Test Kit | - | 1,000 | 14-Mar-2024 | 29-Mar-2024 |

**Product Specification for Merilisa Dengue IgM + IgG Antibody Detection Rapid Test Kit**

| विनिर्देश\|Specification | उप-विनिर्देश\|Sub-Spec | मूल्य\|Value |
|---|---|---|
| GENERAL FEATURES | Product Description | Dengue Rapid Test Kit |
| | Clinical Purpose | To diagnose dengue virus infection in a patient's blood sample |
| PRODUCT INFORMATION | Type of Kit | Dengue IgM + IgG Antibody Detection Rapid Test Kit |
| | Detection Type | Qualitative |
| | Kit should be able to detect all the 4 serotypes of dengue viruses (DEN-1, DEN-2, DEN-3, and DEN-4) | Yes |
| | The test should be able to differentially detect IgG and IgM Antibodies against all 4 serotypes of Dengue virus | Yes |
| | Test should have no cross reactivity with other Flavivirus group mediated and mosquitoes-borne disease | Yes |
| | Testing Principle | Lateral Flow Immunochromatographic Assay |
| | Species Reactivity | Human |
| | Type of Sample | Serum,Plasma |
| | Time to Result | ≤ 30 minutes |
| | Ability to evaluate negative or positive test result | Yes |
| | Sensitivity for Dengue NS1 Ag (%) | NA for IgM/IgG only kit |
| | Specificity for Dengue NS1 Ag (%) | NA for IgM/IgG only kit |
| | Sensitivity for Dengue IgM/IgG Antibody (%) | ≥ 94% |
| | Specificity for Dengue IgM/IgG Antibody (%) | ≥ 96% |
| | Declared sensitivity and specificy 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 | 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 |
| PACKAGING | Pack size of kit | 10 Tests |
| | The test kit packed in such a way that there is proivision 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 made there under as amended till date | Yes |
| | Valid Drug License Number | MFG/MD/2018/000029 |
| | Manufacturing unit certification | ISO:13485 (Latest) |
| | Additional voluntary certification available | GMP/1807943/2018 |
| | 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 | 18 |
| | 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 | No |

| | | |
|---|---|---|
| | bidding | |
| 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 SPECIAL TERMS AND CONDITIONS (STC)**

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**