Bid Number: GEM/2020/B/705562
Dated: 15-07-2020

# Bid Document

| Bid Details | |
|---|---|
| Bid End Date/Time | 18-07-2020 17:00:00 |
| Bid Opening Date/Time | 18-07-2020 17:30:00 |
| Bid Life Cycle (From Publish Date) | 90 (Days) |
| Bid Offer Validity (From End Date) | 30 (Days) |
| Ministry/State Name | Uttar Pradesh |
| Department Name | Medical Health And Family Welfare Department Uttar Pradesh |
| Organisation Name | N/a |
| Office Name | Lucknow Hq |
| Total Quantity | 200000 |
| Item Category | Rapid Antigen Test Kits for Novel Corona Virus (SARS-CoV-2/COVID-19) |
| Bidder Turnover (Last 3 Years) | 300 Lakh (s) |
| MSE Exemption for Years of Experience and Turnover | No |
| Startup Exemption for Years of Experience and Turnover | No |
| Document required from seller | Bidder Turnover,Certificate (Requested in ATC),OEM Authorization Certificate<br>*In case any bidder is seeking exemption from Experience / Turnover Criteria, the supporting documents to prove his eligibility for exemption must be uploaded for evaluation by the buyer |
| Bid to RA enabled | No |
| Inspection Required | No |
| Estimated Bid Value | 100800000 |

**EMD Detail**

| | |
|---|---|
| Advisory Bank | State Bank of India |
| EMD Percentage(%) | 2.00 |
| EMD Amount | 2016000 |

**ePBG Detail**

| | |
|---|---|

| | |
|---|---|
| Advisory Bank | State Bank of India |
| ePBG Percentage(%) | 5.00 |
| Duration of ePBG required (Months). | 14 |

(a). EMD EXEMPTION: The bidder seeking EMD exemption, must submit the valid supporting document for the relevant category as per GeM GTC with the bid. Under MSE category, only manufacturers for goods and Service Providers for Services are eligible for exemption from EMD. Traders are excluded from the purview of this Policy.

(b). EMD & Performance security should be in favour of Beneficiary, wherever it is applicable.

**Beneficiary:**
Managing Director
Lucknow
(Shruti Singh)

**Splitting**

Bid splitting not applied.

## Rapid Antigen Test Kits For Novel Corona Virus (SARS-CoV-2/COVID-19) ( 200000 Test )

**Technical Specifications**

* As per GeM Category Specification

| Specification | Specification Name | Values | Bid Requirement (Allowed Values) |
|---|---|---|---|
| GENERAL FEATURES | Type of Kit | Rapid Antigen Test Kit for Novel Coronavirus (SARS-CoV-2/COVID-19) | * |
| | Purpose | For in-vitro diagnosis/detection of novel coronavirus (SARS-CoV-2/COVID-19) infection | * |
| PRODUCT INFORMATION | Detects | Antigen specific for SARS-CoV-2 | * |
| | Detection Method | Qualitative | * |
| | Testing Principle | Lateral flow chromatographic immunoassay | * |
| | Species Reactivity | Human | * |

| | | | |
|---|---|---|---|
| | Test can be performed using | Nasopharyngeal swab | * |
| | Result reading time | With in 10 to 30 minutes | * |
| | Ability to evaluate negative or positive test result | Yes | * |
| | Contains an internal control Line/dot/band for the confirmation that the test has been performed correctly | Yes | * |
| | Assay Sensitivity | 84 | * |
| | Assay Specificity | 100 | * |
| | Declared sensitivity and specificity shall be claimed by the manufacturer in the kit literature | Yes | * |
| | No cross reactivity | Yes | * |
| | Storage temperature of Kit | 2 to 30° C | * |
| | Shipping Condition | At Room Temperature | * |
| KIT CONTENTS | Kit contents per test | 1 Test cassette with desiccant, 1 sterile swab, 1 Extraction tube with buffer (if any) and 1 nozzle cap/dropper | * |
| | Type of Swab provided | Nasopharyngeal swab | * |
| | Material of Swab | Synthetic fibre swab (nylon, polyester, rayon, or dacron) with plastic shafts or wire shaft (flexible shaft) | * |
| | All the components shall be in the quantity as per pack size | Yes | * |
| | Adequate document in English detailing principle, components, methodologies, validity criteria, interpretation of result, performance characteristic, bio-safety, limitation of assay, storage condition, mfg and exp date & method of disposal provided | Yes | * |
| | | | |

| | | | |
|---|---|---|---|
| | Positive and negative controls provided with each pack of kit | No | * |
| INTERPRETATION OF RESULTS | Reading of results | Visually | * |
| | For all the samples tested, the control line shall be visible at the end of the test | Yes | * |
| | Test will be regarded as invalid if the control line is not visible at the end of the test | Yes | * |
| | Test will be regarded as positive If test line is visible along with the control line | Yes | * |
| PACKAGING | Pack size of the kit | 25 Tests | * |
| | The test kit packed in such a way that there is proivision to conduct single test at a time | Yes | * |
| | Packing of test cassette | Each test cassette with desiccant individually packed in a hermetically sealed and non-permeable pouch | * |
| | Packing of Swab | Each swab individually packed in easy to open peel pouches | * |
| ADDITIONAL REQUIREMENT | Additional Requirement | NA | * |
| CERTIFICATIONS & REPORTS | The kit approved from the statutory authority in its country of origin | Yes | * |
| | Imported kits registered and licensed in India by DCGI & Indigenous manufactured kits licensed by the authority defined under Drugs and Cosmetics act 1940 and Medical Devices Rules, 2017 as amended till date | Yes | * |
| | Availability of certificate/License from the Drugs Controller General of India (DCGI) for import (in case of | Yes | * |

| | | | |
|---|---|---|---|
| | imported kits) and marketing of the intended item | | |
| | Availability of valid drug license for the product issued from the competent authority defined under Drugs & Cosmetics Act, 1940 and Medical Devices Rules, 2017 as amended till date | Yes | * |
| | Drug License Number | - | * |
| | Product certifications/approvals | Kit evaluated and validated by ICMR-NIV-Pune or any other ICMR validation centres | * |
| | Manufacturer facility certifications | ISO:13485 (Latest) | * |
| | Availability of Test report of each batch of the product to prove the conformity to the declared specification | In house Test report from the manufacturer | * |
| | Availability of validation report | Yes | * |
| | Submission of all required certifications, license , test reports and any other specific certification requirement as imposed by country of origin to the buyer at the time of bidding in case of bidding/ at the time of supply in case of direct order | Yes | * |
| SHELF LIFE | **Shelf life from the date of manufacture** | 24 | 12, 18, 24 Or higher |
| | Minimum shelf life of the product at the time of delivery to the consignee | 3/4 of the 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 | No | * |

* Specifications highlighted in bold are the Golden Parameters.

* Bidders may note that In respect of non-golden Parameters, the specifications 'Values' chosen by Buyer will generally be preferred over 'Bid requirement ( allowed Values) by the Buyer.

**Additional Specification Documents**

---

**Consignees/Reporting Officer and Quantity**

| S.No. | Consignee/Reporting Officer | Address | Quantity | Delivery Days |
|---|---|---|---|---|
| 1 | Shariq Ul Islam | 226010,SUDA Bhawan, 7/23, Sector-7, Gomtinagar Extension, Lucknow-226010 | 200000 | 5 |

# Bid Specific Additional Terms and Conditions

1. **Staggered Delivery:** The ordered items shall be supplied in a staggered manner. ( 100000 Quantity shall be supplied within 3 days of contract placement and thereafter 100000 Quantity per Week)
2. OPTION CLAUSE: The Purchaser reserves the right to increase or decrease the quantity to be ordered up to 25 percent at the time of placement of contract. The purchaser also reserves the right to increase the ordered quantity by up to 25% of the contracted quantity during the currency of the contract at the contracted rates. Bidders are bound to accept the orders accordingly.
3. Scope of supply (Bid price to include all cost components) : Supply Installation Testing Commissioning of Goods and Training of operators and providing Statutory Clearances required (if any)
4. The bidder is required to upload, along with the bid, all relevant certificates such as BIS licence, type test certificate, approval certificates and other certificates as prescribed in the Product Specification given in the bid document.
5. Bidder Turn Over Criteria: The minimum average annual financial turnover of the bidder during the last three years, ending on 31st March of the previous financial year, should be as indicated in the bid document. Documentary evidence in the form of certified Audited Balance Sheets of relevant periods or a certificate from the Chartered Accountant / Cost Accountant indicating the turnover details for the relevant period shall be uploaded with the bid. In case the date of constitution / incorporation of the bidder is less than 3 year old, the average turnover in respect of the completed financial years after the date of constitution shall be taken into account for this criteria.

This Bid is also governed by the General Terms and Conditions

**---Thank You---**