

Bid Document

| Bid Details | |
|--|---|
| Bid End Date/Time | 05-08-2021 16:00:00 |
| Bid Opening Date/Time | 05-08-2021 16:30:00 |
| Bid Life Cycle (From Publish Date) | 90 (Days) |
| Bid Offer Validity (From End Date) | 70 (Days) |
| Ministry/State Name | Chandigarh |
| Department Name | Education Department Chandigarh |
| Organisation Name | Government Medical College And Hospital |
| Office Name | Sector 32, Chandigarh |
| Total Quantity | 300 |
| Item Category | sutures |
| MSE Exemption for Years of Experience and Turnover | No |
| Startup Exemption for Years of Experience and Turnover | No |
| Bid to RA enabled | No |
| Time allowed for Technical Clarifications during technical evaluation | 2 Days |
| Evaluation Method | Total value wise evaluation |

EMD Detail

| | |
|-------------------|---------------------|
| Advisory Bank | State Bank of India |
| EMD Percentage(%) | 2.00 |
| EMD Amount | 25671 |

ePBG Detail

| | |
|----------|----|
| Required | No |
|----------|----|

(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:

Director Principal
Government Medical College and Hospital, Sector 32 ,Chandigarh
(Jasbinder Kaur)

Splitting

Bid splitting not applied.

MSE Purchase Preference

| | |
|-------------------------|----|
| MSE Purchase Preference | No |
|-------------------------|----|

MII Purchase Preference

| | |
|-------------------------|----|
| MII Purchase Preference | No |
|-------------------------|----|

Sutures (300 box)

| | |
|------------|------------------|
| Brand Type | Registered Brand |
|------------|------------------|

Technical Specifications

[* As per GeM Category Specification](#)

| Specification | Specification Name | Bid Requirement (Allowed Values) |
|-------------------------|--|----------------------------------|
| Performance Parameters | Governing Specification | USP |
| | category of suture | absorbable |
| | Construction | Braided |
| | Suture Material | Polyglactin |
| | Suture Length (cm) | 90 |
| | Suture Size | 2-0 |
| | Needle | With |
| Needle Parameters | Type of Needles | Round bodied |
| | Needle Length (mm) | 30 |
| | Needle Curvature | 1/2 circle |
| Packing Parameters | Number of suture foils in box | 12 |
| Certification & Reports | Availability of any other certification such as CE/FDA/CSA/PQS /ISO etc... | Yes |

Consignees/Reporting Officer and Quantity

| S.No. | Consignee/Reporting Officer | Address | Quantity | Delivery Days |
|-------|-----------------------------|---|----------|---------------|
| 1 | Sushil Kumar Yadav | 160031, Government Medical College & Hospital Sector 32, Chandigarh, 160031 | 300 | 30 |

Special terms and conditions-Version:1 effective from 04-05-2020 for category sutures

- Special Terms and Conditions for Medical Devices and Consumables covered under Provisions of Drug and Cosmetic Act
 - For items wherever Drug Licence requirements are applicable all provisions of Drug and Cosmetic Act 1940 as amended up to date and Rules made there under will be applicable in addition to any other terms and conditions specified in the Portal.
 - Drug License: For indigenous products offered in the market, Manufacturer should have valid Drug License as per Drugs and Cosmetic Act 1940 issued by concerned State Drug Control authorities .The Seller if different from the manufacturer shall also be required to be holding Drug License for sale . In case of imported products Manufacturer shall be registered under Form no 10 with Central Drug Authorities (CDSCO) and the Seller offering imported products should be also holding valid Sales License issued by the local drug authorities. For imported products, certificate from the OEM that product is being used in the Country of Origin should be available with the Seller. It shall be the responsibility of the Seller to ensure that that the Drug License is valid for the product offered and due to any reason the drug control authorities have cancelled or suspended Drug License or convicted the manufacturer or Seller for any offence under the provisions of Drug and Cosmetic Act, Seller should immediately withdraw the product and also intimate the Buyers in case of pending orders for supplies as well as the GeM administration regarding the matter.
 - Manufacturing & Marketing Experience: Sellers offering the Products in the Portal either as Manufacturers or as Authorised Seller shall ensure that the Products offered are being Manufactured and Marketed in the country (for Indigenous Products) and Marketed (for Imported Products) continuously at least for the last 2 years
 - Certifications: Manufacturers of offered product (Offered by Manufacturers or by Authorized Seller) should be holding valid Good Manufacturing Practices Certificate (GMP) as per revised Schedule-M of Drug and Cosmetic Act 1940 as amended up to date or WHO-GMP as per norms amended up to date issued by the Licensing Authority or certificate which is at par with WHO-GMP issued by the authorities of exporting countries / COPP certificate .
 - Non Conviction Certificate: Sellers either Manufacturers or Authorized Sellers are required to ensure that they are not under conviction in terms of the provisions of Drugs & Cosmetic Act and any other law applicable in relation to the same . In case at any point of time, the Manufacturer or Authorized Seller is convicted under provisions of Drug and Cosmetic Act, it shall be their responsibility to withdraw the product immediately from the market.
 - Banning and Blacklisting: Seller either Manufacturer or Authorized Seller shall ensure that there is no banning or black listing applicable against them for the product offered on the portal due to quality failure and /or fraudulent/illegal practices or for any other reasons
 - It shall be the responsibility of the Seller either Manufacturer or Authorized Seller to ensure that manufacturer is having own in-house testing lab to carry out all the required tests as per specification and provisions of drug act as amended up to date for the quoted product and shall also forward the copies of the in-house test reports for each batch along with the supplies. For imported products , certificate from OEM regarding availability of all test facilities in house with them should be available with Seller .
 - Each lot of supplies shall be dispatched under Self Certification scheme duly supported by in house test reports. Consignees shall be at liberty to draw control Samples and send it to approved Laboratories for testing and in case of any failure , entire responsibility shall rest with Seller in addition to any penalties under the provisions of Drug Act including removal of Goods from the Consignee place. .Further administrative actions as per terms and conditions Gem Portal shall also be applicable.
 - Packing shall be as per relevant clause of Standard Specifications applicable as indicated in the Catalogue Parameters indicated in the Portal and as per provisions of Drug & Cosmetic Act as amended up to date.
 - Marking: Each Primary Packing shall be marked as under:-
 - Nomenclature of the stores

2. Manufacturers Name, Address, Drug License No.
3. Month of manufacturing, Expiry, Batch No and lot No (if applicable)
4. Any other particulars required under Drug and Cosmetic Act 1940 amended up to date if item is governed under drug and cosmetic act
5. Quantity contained therein
6. Manufacturers Name or Trade Mark
7. Government Supply - ""Not For Sale
8. Secondary Packing Cartons shall be marked with Manufacturers Name, Batch no and Month of Manufacture and Use Before.

Buyer Added Bid Specific Additional Terms and Conditions

1. Scope of supply (Bid price to include all cost components) : Only supply of Goods

Disclaimer

The additional terms and conditions have been incorporated by the Buyer after approval of the Competent Authority in Buyer Organization. Buyer organization is solely responsible for the impact of these clauses on the bidding process, its outcome and consequences thereof including any eccentricity / restriction arising in the bidding process due to these ATCs and due to modification of technical specification and / or terms and conditions governing the bid. Any clause incorporated by the Buyer such as demanding Tender Sample, incorporating any clause against the MSME policy and Preference to make in India Policy, mandating any Brand names or Foreign Certification, changing the default time period for Acceptance of material or payment timeline governed by OM of Department of Expenditure shall be null and void and would not be considered part of bid. Further any reference of conditions published on any external site or reference to external documents / clauses shall also be null and void. If any seller has any objection / grievance against these additional clauses or otherwise on any aspect of this bid, they can raise their representation against the same by using the Representation window provided in the bid details field in Seller dashboard after logging in as a seller within 4 days of bid publication on GeM. Buyer is duty bound to reply to all such representations and would not be allowed to open bids if he fails to reply to such representations.

[This Bid is also governed by the General Terms and Conditions.](#)

In terms of GeM GTC clause 26 regarding Restrictions on procurement from a bidder of a country which shares a land border with India, any bidder from a country which shares a land border with India will be eligible to bid in this tender only if the bidder is registered with the Competent Authority. While participating in bid, Bidder has to undertake compliance of this and any false declaration and non-compliance of this would be a ground for immediate termination of the contract and further legal action in accordance with the laws.

---Thank You---