



JAMMU AND KASHMIR MEDICAL SUPPLIES CORPORATION LTD.

(Public Sector Undertaking of Govt of Jammu & Kashmir)

Corporate Head Office: 1st Floor, Drug Store Building, Govt. Medical College, Bakshi Nagar, Jammu

Temporary shifted to Satyam Road, Plot No. 58 Friends colony Trikuta Nagar Jammu

Corporate Office Kashmir: Opposite State Motor Garage Deptt. Baramulla National Highway, Bemina (190018) Srinagar

Telephone: 0191-2580842, Fax: 0191-2581845 (Jammu)

MINOR INFIRMITIES/SHORTCOMINGS NOTICE-502

"Human Factor-VIII with VWF Concentrate"

Subject: Evaluation of Technical bids for finalization of the contract for the Procurement of "Human Factor-VIII with VWF Concentrate".

Report of the Sub-Committee constituted for evaluation of bid documents for the finalization of the rate contract for the procurement of "Human Factor-VIII with VWF Concentrate" with reference to the NIT No: JKMSCL/Human Factor-VIII with VWF Concentrate/2021/502 dated. 20.11.2021. The preliminary evaluation of the bid documents have been done on the basis of uploaded documents.

Documents of the following bidders who participated in the tender were evaluated and the shortcomings found in the documents are shown against each:

S. No	Name of the bidder	Minor Infirmitities/Shortcomings
1/4	M/S Alpha Drugs, New Delhi. (Bidder/Importer) M/S Bio Products Laboratory Ltd, UK (Manufacturer) Brand Name: Optivate®	<ol style="list-style-type: none">1. Average Annual Turnover Statement not less than 20 crores of the Importer (M/S Alpha Drugs, New Delhi) for last 3 financial years from Chartered Accountant with UDIN (2017-18, 2018-19 & 2019-20). (Submitted document is of Foreign Manufacturer.)2. Copies of Audited Balance Sheet & Profit Loss Account of the Importer (M/S Alpha Drugs, New Delhi) for last 3 financial years from Chartered Accountant (2017-18, 2018-19 & 2019-20).3. Submit valid Drug Manufacturing License along with subsequent renewals of Original Manufacturer(s) for the quoted product.4. Submit valid CGMP as per revised Schedule "M"/WHO format of the manufacturer for the quoted item. (Submitted document is not for the quoted item)5. Product permission by the licensing authority for the quoted products by the Original Manufacturer to be submitted.6. Non conviction certificate of the Principal Manufacturer (M/S Bio Products Laboratory Ltd.) for quoted product from licensing authority for the Import license no. BP-7 to be submitted.7. Market standing certificate issued by licensing authority of the respective state not less than three preceding year.(2017-18, 2018-19 & 2019-20) to be submitted . (Submitted is for Anti-Haemophillie Factor VIII not for Human Factor VIII with vWF concentrate.)
2/4	M/S Grifols India Healthcare (P) Ltd, Haryana (Bidder/Importer)	<ol style="list-style-type: none">1. Submit valid CGMP as per revised Schedule "M"/WHO format of the manufacturer for the quoted item.


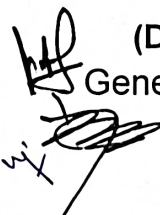
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	<p>M/S Grifols Biologicals LLC, USA (Manufacturer)</p> <p>Brand Name: Alphanate.</p>	<ol style="list-style-type: none"> Product permission by the licensing authority for the quoted products by the Original Manufacturer to be submitted. Authorization letter from principal manufacturer in favour Importer (M/S Grifols India Healthcare (P) Ltd, Haryana) for the quoted product to be submitted. Market standing certificate issued by licensing authority of the respective state not less than three preceding year.(2017-18, 2018-19 &2019-20) to be submitted for the quoted product Human Factor VIII with vWF concentrate .
3/4	<p>M/S Intas Pharmaceuticals Ltd., Gujarat. (Bidder/manufacturer)</p> <p>M/s Novolife Healthcare Corp., Srinagar. (Authorized Representative)</p> <p>Brand Name:</p>	<ol style="list-style-type: none"> Submit valid Drug Manufacturing License along with subsequent renewals of Original Manufacturer(s) for the quoted product as per NIT Specification. (The submitted document mentions Dried Human Anti-Haemophilic Factor VIII and does not Mentions vWF as per NIT specification.) Submit valid CGMP as per revised Schedule "M"/WHO format of the manufacturer for the quoted item. (The submitted document mentions Dried Human Anti-Haemophilic Factor VIII and does not Mentions vWF as per NIT specification.) Product permission by the licensing authority for the quoted products by the Original Manufacturer to be submitted. (The submitted document mentions Dried Human Anti-Haemophilic Factor VIII and does not Mentions vWF as per NIT specification.) Market standing certificate issued by licensing authority of the respective state not less than three preceding year.(2017-18, 2018-19 &2019-20) to be submitted for the quoted product Human Factor VIII with vWF concentrate .(The submitted document mentions Dried Human Anti-Haemophilic Factor VIII and does not Mentions vWF as per NIT specification.) Valid Drug sale License along with subsequent renewals on form 20B & 21B to be submitted of M/s Novolife Healthcare Corp., Srinagar.(Authorized Representative) Latest Non-Conviction certificate issued by licensing authority to be submitted of M/s Novolife Healthcare Corp., Srinagar (Authorized Representative) issued not before 06 months. Clarify the Brand Name of the product Quoted.
4/4	<p>M/s Baxalta Bioscience India Pvt. Ltd., Gurugram. (Bidder/Importer)</p> <p>M/s Baxter AG., Austria. (Manufacturer)</p> <p>Brand Name: Immunate.</p>	<ol style="list-style-type: none"> Submit valid Drug Manufacturing License along with subsequent renewals of Original Manufacturer(s) for the quoted product as per NIT Specification. (The submitted document mentions Purified freeze Dried Human coagulation Factor VIII and does not Human Factor VIII with vWF concentrate as per NIT specification.) Submit valid CGMP as per revised Schedule "M"/WHO format of the manufacturer for the quoted item. (The submitted document mentions Purified freeze Dried Human coagulation Factor VIII and does not Human Factor VIII with vWF concentrate as per NIT specification.) Product permission by the licensing authority for the quoted

		<p>products by the Original Manufacturer to be submitted. (The submitted document mentions Purified freeze Dried Human coagulation Factor VIII and does not Human Factor VIII with vWF concentrate as per NIT specification.)</p> <p>4. Market standing certificate issued by licensing authority of the respective state not less than three preceding year.(2017-18, 2018-19 &2019-20) to be submitted for the quoted product Human Factor VIII with vWF concentrate .(The submitted document mentions Purified freeze Dried Human coagulation Factor VIII and does not Human Factor VIII with vWF concentrate as per NIT specification.)</p> <p>5. Import license on Form 40 & registration approved by CDSCO.</p> <p>6. List of items (Annexure C) highlighting the items quoted Human Factor VIII with vWF concentrate by the bidder mentioning the principal manufacturer of quoted item and brand name to be submitted. .(The submitted document mentions Purified freeze Dried Human coagulation Factor VIII and does not Human Factor VIII with vWF concentrate as per NIT specification.)</p>
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The bidders are requested to upload the infirmities by or before **14-01-2022 up to 4:00 P.M** on gmjkmscl.mi@gmail.com which shall be forwarded to technical evaluation Sub-Committee for further scrutiny and final recommendation in the matter.

In case the documents are not submitted on specified date and time ,your bid shall be evaluated only on the basis of documents submitted earlier, documents received after due date and time shall not be considered and no hard copy of the documents will be entertained.


 (Dr. Ab. Majid Mirab)
 General Manager (Drugs)
 JKMSCL


No: JKMSCL/GM (Drugs)/NIT-502/M.I/2022/10622-23

Dated: 08-01-2022

Copy for information to the:-

1. P.A to Managing Director, JKMSCL
2. Asstt. Programmer (M) JKMSCL I/c website to uploaded the Minor Infirmities on the official/relevant website.
3. Office file.