

JAMMU AND KASHMIR MEDICAL SUPPLIES CORPORATION LTD.

(Public Sector Undertaking of Govt of Jammu & Kashmir)

Corporate Head Office: Plot No: 58, Friends Colony, Satyam Road, Trikuta Nagar Jammu Corporate Office: Opp State Motor Garages Near Haj House, Bemina, Srinagar Telephone: 0191-2580842, Fax: 0191-2581845 (Jammu); Telefax: 0194-2432008 (Srinagar)

NOTICE

Sub:

Shortcomings /minor infirmities of tenders invited for finalization of the contract for the procurement of Blood Bank Items*

Ref:

. JKMSCL/MED /2021/ 449 DATED: 18-01-2021

The evaluation of technical documents against aforesaid NIT was done by the Sub-Committee. Some documents from the participating bidders, are found deficit as per the Notice Inviting Bid. With reference to clause 2.1.9- Chapter II of Standard Procurement Procedure approved during 2nd Board Meeting, the under mentioned shortcomings comes under minor infirmities. It is therefore, impressed upon the under mentioned firms to upload the shortcomings/ clarifications asked for, against each, latest by 09-08-2021 upto 1300 hrs., failing which the e-bids of the respective firms shall be liable to be rejected. The desired documents may be submitted on mail id:gmjkmscl.mi@gmail.com or can be uploaded on the web site of JKMSCL www.jkmsclbusiness.com.

S. No	Name of the bidder/ manufacturer	Item quoted	Documents (infirmities) required
1	M/s Navdeep Sales Con Jammu(Bidder)	rporation Ltd,	Latest GST Return of the bidder to be submitted.
	M/s Voxture Bio Ltd. Maharashtra (Manufacturer)	BB-01, Anti Sera A BB-02, Antisera-B BB-04, Anti Sera D (Blend) BB-06 Anti AB	Copies of Audited Balance Sheet & Profit Loss Account for last three financial years from Chartered Accountant to be submitted.
		BB-07, Detection for malarial parasite BB-08, Detection for	Details of Technical personnel employed in the manufacturing and testing approved by the Licensing Authority to be submitted
		Syphilis VDRL Kit BB-020, HIV Elisa Testing Kit IV generation	BIS License with schedule for ISI Marked Products Quoted (wherever applicable) to be submitted.
		BB-022, Hepatitis B Surface Antigen testing	Product permission for Hepatitis-B; BB- 022 to be submitted.
		Kits	Market standing certificate issued by licensing authority of the respective states not less than three preceding years to be submitted. Submitted is not notarized and valid till 14-05-20.
	M/s Mitra Industries Pvt. Ltd. Faridabad (Manufacturer)	BB-09, Top & Bottom blood bag or top and top for collection of 350 ml & 450 ml of whole blood with integral leukocyte filter for red cell filtration. BB-10, Single Blood Bags Capacity: 450ml BB-11, Single Blood Bags Capacity: 350ml BB-012, Double blood bag (350ml) Double Bag Primary bag (350ml).	Copies of Audited Balance Sheet & Profit Loss Account for last three financial years from Chartered Accountant to be submitted. Details of Technical personnel employed in manufacturing and testing approved by the Licensing Authority to be submitted. BIS License with schedule for ISI Marked Products Quoted (wherever applicable) to be submitted.



		Bag(450ml) with SAGI BB-16, Quadruple Blag (350ml) (w SAGM): Capic Quadruple blood bag BB-17, Quadruple Blag (450ml) (with SAGI	lood Bag ml). Dml) Bag ood M ood with city:
2	M/s Novex Life Sciences, Jammu (Bidder)	Authorization from pri Delux Surgical to be si	ncipal manufacturer / Importer from M/s
	M/s Levram Lifesciences Pvt. Ltd. Silvasaa (Manufacturer)		
	M/s Delux Surgicals, Ahmadabad (Manufacturer)	BB-018, Blood Transfusion set	Copies of Audited Balance Sheet & Profit Loss Account for last three financial years from Chartered Accountant is not legible Valid CGMP as per revised Schedule "M"/ WHO format to be submitted Statement of Installed Manufacturing
			Capacity, Certificate regarding rate reasonability and details of warehouses (Annexure -E) to be submitted



Non blacklisting / undertaking declaration(notarized) (Original manufacturer/ Direct Importer) on nonjudicial Stamp paper of Rs 100/- (Annexure-F) to be submitted Market Standing Certificate issued by Licensing authority of the respective States not Less than Three preceding years. (duly notarized) Annexure G to be submitted Details of Technical personnel employed in the manufacturing and testing approved by the Licensing Authority to be submitted. BIS License with schedule for ISI Marked Products Quoted (wherever applicable) to be submitted Product permission of Blood Transfusion Set to be submitted Submitted Average Annual Turnover is not readable. Submit readable copy of Average Annual turnover statement. Authorization from principal manufacturer in favour of bidder to be submitted. M/s Diagast India, BB-01, Anti Sera Copies of Audited Balance Sheet & Profit Authorized representative Loss Account for last three financial years of M/s Diagast France from Chartered Accountant to be submitted. (manufacturer) BB-02, Antisera-Valid Manufacturing License along with subsequent renewals, if any to be submitted BB-03, Anti Sera Product permission by the licensing authority for the quoted products by the original BB-04, Anti Sera manufacturer(s) to be submitted D (Blend) Statement of Installed Manufacturing Capacity, Certificate regarding rate reasonability and details of warehouses (Annexure -E) to be submitted Non blacklisting / undertaking declaration(notarized) (Original manufacturer/ Direct Importer) on nonjudicial Stamp paper of Rs 100/- (Annexure-F) to be submitted Market Standing Certificate issued by Licensing authority of the respective States not Less than Three preceding years. (duly notarized) Annexure G to be submitted Details of Technical personnel employed in the manufacturing and testing approved by the Licensing Authority to be submitted BIS License with schedule for ISI Marked Products Quoted (wherever applicable) to be submitted ISO & CE/BIS/USFDA certificate for quoted Items as mentioned in bid Catalogue (wherever applicable) to be submitted. Average Annual Turnover statement not less than 1 crores (One Crore) of the bidder (original manufacturer/importer) for last 3 financial years from Chartered Accountant with UDIN to be submitted.

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			Valid CGMP as per revised Schedule- M/WHO format to be submitted. Authorization of M/s Diagast Frances in favour of Diagast India to be submitted.
	M/s Trivitron Healthcare Pvt. LTd. Tamil Nadu (Manufacturer) BB-020, HIV Elisa Testing Kit IV generation BB-021, HCV Elisa Testing Kit IV generation. BB-022, Hepatitis B Surface Antigen testing Kits	Elisa Testing Kit	Valid Manufacturing License along with subsequent renewals, if any to be submitted
		Licensing authority of the respective States not Less than Three preceding years. (duly notarized) Annexure G to be submitted Details of Technical personnel employed in the manufacturing and testing approved by the Licensing Authority to be submitted. BIS License with schedule for ISI Marked Products Quoted (wherever applicable) to be submitted ISO & CE/BIS/USFDA certificate for quoted Items as mentioned in bid Catalogue (wherever applicable) to be submitted Non blacklisting / undertaking declaration (notarized) (Original manufacturer/ Direct Importer) on non- judicial Stamp paper of Rs 100/-	
3	M/s SR Technomed, J M/s Maco Pharma, New Delhi (Importer)	ammu (Bidder) BB-09, Top & Bottom blood bag or top and	Average Annual Turnover Statement not less than 1 crores (One crore) of the bidder/ Importer for Last 3 financial Years from Chartered Accountant with UDIN ANNEXURE-F to be submitted
	M/s Macopharma India Transfusion	top for collection of 350 ml & 450 ml of whole blood with integral	
	Solution Pvt. Ltd. New Delhi (Factory at Poland & France)	leukocyte filter for red cell filtration. BB-17, Quadruple	Latest Non Conviction Certificate issued by the Licensing authority of the respective state / UT (Issued not before 6 months) to be submitted.
	M/s Maco Pharma France, Poland (Manufacturer)	Blood Bag (450ml) (with SAGM)	Copies of Audited Balance Sheet & Profit Loss Account for last three financial years from Chartered Accountant to be submitted
			Valid Manufacturing License along with subsequent renewals, if any to be submitted
			Statement of Installed Manufacturing Capacity, Certificate regarding rate reasonability and details of warehouses (Annexure –E) to be submitted
			Valid CGMP as per revised Schedule "M"/ WHO format to be submitted Non blacklisting / undertaking
	The state of the s		declaration(notarized) (Original manufacturer/ Direct Importer) on non-judicial Stamp paper of Rs 100/- (Annexure-F) to be submitted



		Market Standing Certificate issued by Licensing authority of the respective States not Less than Three preceding years. (duly notarized) Annexure G to be submitted BIS License with schedule for ISI Marked Products Quoted (wherever applicable) to be submitted. Authorization from foreign manufacturer in favour of important to be submitted.
M/s Meril Diagnostis Pvt. Ltd. (Mumbai) (manufacturer)	BB-01, Anti Sera A BB-02, Antisera-B BB-03, Anti Sera D BB-04, Anti Sera D (Blend) BB-05m / Anti-A1 lectin BB-06 Anti AB BB-07, Detection for malarial parasite BB-08, Detection for Syphilis VDRL Kit BB-020, HIV Elisa Testing Kit IV generation BB-021, HCV Elisa Testing Kit IV generation. BB-022, Hepatitis B Surface Antigen testing Kits	favour of importer to be submitted. Copies of Audited Balance Sheet & Profit Loss Account for last three financial years from Chartered Accountant to be submitted Valid CGMP as per revised Schedule "M"/WHO format to be submitted Latest Non Conviction Certificate issued by the Licensing authority of the respective state / UT (Issued not before 6 months) to be submitted Product permission by the licensing authority for the quoted products by the original manufacturer(s) to be submitted Valid Manufacturing License along with subsequent renewals, if any to be submitted ISO & CE/BIS/USFDA certificate for quoted Items as mentioned in bid Catalogue (wherever applicable) to be submitted Non blacklisting / undertaking declaration(notarized) (Original manufacturer/Direct Importer) on non-judicial Stamp paper of Rs 100/- (Annexure-F) to be submitted Statement of Installed Manufacturing Capacity, Certificate regarding rate reasonability and details of warehouses (Annexure -E) to be submitted Market Standing Certificate issued by Licensing authority of the respective States not Less than Three preceding years. (duly notarized) Annexure G to be submitted Details of Technical personnel employed in the manufacturing and testing approved by the Licensing Authority to be submitted BIS License with schedule for ISI Marked Products Quoted (wherever applicable) to be submitted
M/s Becton Dickinson India Pvt.Ltd. Gurgaon (Importer) M/s Becton Dickinson & Factory at USA, UK, India (Manufacturer)	BB-019, K2EDTA Vacutainer	Average Annual Turnover Statement not less than 1 crores (One crore) of the bidder (Original Manufacturer / Importer) for Last 3 financial Years from Chartered Accountant with UDIN ANNEXURE-F to be submitted. Valid CGMP as per revised Schedule "M"/WHO format to be submitted. Product permission by the licensing authority for the quoted products by the original manufacturer(s) to be submitted. Statement of Installed Manufacturing Capacity, Certificate regarding rate reasonability and details of warehouses (Annexure -E) to be submitted.

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				Valid Manufacturing License along with subsequent renewals, if any to be submitted.	
				Non blacklisting / undertaking declaration (notarized) (Original manufacturer/ Direct Importer) on non-judicial Stamp paper of Rs 100/- (Annexure-F) to be submitted. Details of Technical personnel employed in the manufacturing and testing approved by the Licensing Authority to be submitted.	
				BIS License with schedule for ISI Marked Products Quoted (wherever applicable) to be submitted ISO & CE/BIS/USFDA certificate for quoted Items as mentioned in bid Catalogue (wherever applicable) to be submitted.	
4	Private Ltd, Kerela (Bidder/Manufactur er) Local distributor Jammu Zone: M/s Novex Lifesciences Bakshi Nagar, Jammu Srinagar Zone: M/s Global scientific enterprises, Karan Nagar, Srinagar BB-09, Top & Bottom blood bag or top and top for collection of 350 ml & 450 ml of whole blood with integral leukocyte filter for red cell filtration. BB-012, Double blood bag (350ml) Double Bag Primary bag (350ml). One Satellite bag (300ml) BB-013, Double blood bag (450ml) Double Bag Primary bag (450ml). One Satellite bag (300ml) BB-14, Triple Blood Bag (350ml) with SAGM BB-15, Triple Blood Bag (450ml) with SAGM BB-16, Quadruple Blood Bag (350ml) (with SAGM): Capicity: Quadruple blood bag BB-17, Quadruple Blood Bag (450ml) (with SAGM)		d bag or top and or collection of 350 a 450 ml of whole d with integral ocyte filter for red iltration. 112, Double blood 350ml) Double Primary bag ml). One Satellite 300ml) 113, Double blood 450ml) Double Primary bag ml). One Satellite 300ml) 4, Triple Blood 350ml) with M 5, Triple Blood 350ml) with M 6, Quadruple Bag (350ml) SAGM): Capicity: ruple blood bag 7, Quadruple Bag (450ml)	Details of Technical personnel employed in the manufacturing and testing approved by the Licensing Authority to be submitted BIS License with schedule for ISI Marked Products Quoted (wherever applicable) to be submitted. Product permission by the licensing authority for the quoted products (Blood Bags) to be submitted. Form 20 and 21B of M/s Novex Life Sciences Bakshi Nagar Jammu must be provided to be eligible for point of supply Form 20 and 21B of M/s Global scientific enterprises, Karan Nagar must be provided to be eligible for point of supply	
5	M/s Magnus Enterprises, Jammu (Bidder)		authority of the re months) to be sub		
			to be submitted.	cense along with subsequent renewals, if any	

	M/s Polymedicure Ltd. Haryana (Manufacturer)	bag or to collection of wintegral red cell BB-10, Capacity 450ml BB-11, Capacity 350ml BB-012, (350ml) Primary Satellite BB-14, 7 (350ml) Primary Satellite BB-15, 7 Bag (450 BB-16, Capicity: bag BB-17, CBag (450 BB-18,	op and to on of 350 hole blood leukocyt filtration. Single Bloy: Single Single Bloy: Single y: Single Double B bag (350 bag (300	ml & 450 d with e filter for ood Bags blood bag- ood Bags blood bag- blood bag ag ml). One ml) od Bag ml). One ml) od Bag SAGM e Blood SAGM: ple blood e Blood SAGM):	Product permission by the licensing authority for the quoted products by the original manufacturer(s) to be submitted. BIS License with schedule for ISI Marked Products Quoted (wherever applicable) to be submitted Non blacklisting / undertaking declaration (notarized) (Original manufacturer/ Direct Importer) on non-judicial Stamp paper of Rs 100/-(Annexure-F) to be submitted. Market Standing Certificate issued by Licensing authority of the respective States not Less than Three preceding years. (duly notarized) Annexure G to be submitted.
	M/s Aar Kay Healthcare Pvt. Ltd. Surat, Gujarat (Manufacturer)	BB-02, A BB-03, A BB-04, A BB-05m BB-06 Ar BB-07, D malarial	BB-01, Anti Sera A BB-02, Antisera-B BB-03, Anti Sera D BB-04, Anti Sera D (Blend) BB-05m /Anti-A1 lectin BB-06 Anti AB BB-07, Detection for malarial parasite BB-08, Detection for		Valid CGMP as per revised Schedule "M"/ WHO format to be submitted. BIS License with schedule for ISI Marked Products Quoted (wherever applicable) to be submitted. ISO & CE/BIS/USFDA certificate for quoted Items as mentioned in bid Catalogue (wherever applicable) to be submitted.
6	Irays Meditech Solutions Private Ltd. Gurugram (Bidder)		VDRL Kit Latest author months Name, officer make c	Non Convictity of the rest to be subsphotograph / represents corresponder	tion Certificate issued by the Licensing spective state /UT (Issued not before 6
	M/s Genuine Bio- System, Pvt Ltd. Chennai (Manufacturer)	A A A A A A A A A A A A A A A A A A A		Average As crores (On Last 3 fina	nnual Turnover Statement not less than 1 ne crore) of the Original Manufacturer for ancial Years from Chartered Accountant ANNEXURE-F to be submitted.

		BB-03, Anti Sera	Copies of Audited Balance Sheet & Profit Loss Account for last three financial years from Chartered Accountant to be submitted.	
		BB-04, Anti Sera D (Blend)	Valid Manufacturing License along with subsequen renewals, if any for the quoted items to be submitted.	
		BB-05m /Anti-A1 lectin	manufacturer(s) to be submitted. (Submit documents do not mention the quoted products.	
		BB-06 Anti AB BB-07, Detection for malarial parasite	Non blacklisting / undertaking declaration(notarized) (Original manufacturer/ Direct Importer) on non-judicial Stamp paper of Rs 100/- (Annexure-F) to be submitted. Market Standing Certificate issued by Licensing	
		BB-08, Detection for Syphilis VDRL Kit	authority of the respective States not Less than Three preceding years. (duly notarized) Annexure G to be submitted.	
		All	Details of Technical personnel employed in the manufacturing and testing approved by the Licensing Authority to be submitted	
			Statement of Installed Manufacturing Capacity, Certificate regarding rate reasonability and details of warehouses (Annexure -E) to be submitted. Valid CGMP as per revised Schedule "M"/ WHO format to be submitted.	
			BIS License with schedule for ISI Marked Products	
7	M/s Bhardwaj Agencies, Jammu (Bidder)		Quoted (wherever applicable) to be submitted. Valid Drug sale License along with subsequent renewals, to be submitted. Submitted documents not legible.	
	Most of the docume M/s Bhardwaj Ager readable which neeseriously.	icies are not	Letter of acceptance of Terms and conditions of e- NIT duly signed by the bidder ANNEXURE-L to be submitted.	
	was vowals	·	Latest GST Return of the bidder is not legible. Submitted legible copy of the same	
	M/s Shree Umiya Surgicals Pvt. Ltd. Ahmadabad	BB-018, Blood transfusion set	Valid Manufacturing License along with subsequent renewals, is not legible. Submitted legible copy of the same	
	(Manufacturer)		Valid CGMP as per revised Schedule "M"/ WHO format to be submitted.	
			Product permission by the licensing authority for the quoted products by the original manufacturer(s) is not legible. Submitted legible copy of the same	
			Market Standing Certificate issued by Licensing authority of the respective States not Less than Three preceding years. (duly notarized) Annexure G is not legible. Submitted legible copy of the same	
			Details of Technical personnel employed in the manufacturing and testing approved by the Licensing Authority to be submitted. BIS License with schedule for ISI Marked Products	
			Quoted (wherever applicable) to be submitted. Submit legible copy of the same	
	M/s Avecon Healthgcare Pvt. Ltd, Ambala (Manufacturer)	BB-01, Anti Sera A BB-02, Antisera-	Average Annual Turnover Statement not less than 1 crores (One crore) of the bidder (Original Manufacturer/ Importer) for Last 3 financial Years from Chartered Accountant with UDIN ANNEXURE-	
-		В	F is not legible. Submit legible copy of the same	



	BB-03, Anti Sera D	Copies of Audited Balance Sheet & Profit Loss Account for last three financial years from Chartered Account is not legible. Submit legible copy of the
	BB-04, Anti Sera D (Blend)	Valid Manufacturing License along with subsequent renewals, if any not legible. Submit legible copy of the same
	BB-07, Detection for malarial	Valid CGMP as per revised Schedule "M"/ WHO format to be submitted.
	parasite BB-08, Detection	Product permission by the licensing authority for the quoted products by the original manufacturer(s) to be submitted
	for Syphilis VDRL Kit	Statement of Installed Manufacturing Capacity, Certificate regarding rate reasonability and details o warehouses (Annexure -E) not legible. Submit legible copy of the same
		Market Standing Certificate issued by Licensing authority of the respective States not Less than Three preceding years. (duly notarized) Annexure G to be submitted.
		BIS License with schedule for ISI Marked Products Quoted (wherever applicable) to be submitted.
		Details of Technical personnel employed in the manufacturing and testing approved by the Licensing Authority to be submitted.
		Latest Non Conviction Certificate issued by the Licensing authority of the respective state /UT (Issued not before 6 months) to be submitted.
		Authorization form M/s Avecon Healthcare to be submitted.
		MD-9 of Avecon to be submitted. Submitted copy is not readable.
M/s Bhat Biotech India pvt. Ltd.	BB-06, Anti AB	Valid Manufacturing License along with subsequen renewals, if any to be submitted.
Bangalore (Manufacturer)	BB-020, HIV Elisa Testing Kit IV generation	Statement of Installed Manufacturing Capacity, Certificate regarding rate reasonability and details o warehouses (Annexure –E) is not legible. Submit legible copy of the same
	BB-022, Hepatitis B Surface Antigen testing Kits	Non blacklisting / undertaking declaration(notarized) (Original manufacturer/ Direct Importer) on non-judicial Stamp paper of Rs 100/- (Annexure-F) to be submitted.
		Market Standing Certificate issued by Licensing authority of the respective States not Less than Three preceding years. (duly notarized) Annexure G to be submitted.
		Details of Technical personnel employed in the manufacturing and testing approved by the Licensing Authority to be submitted.
		Product permission by the licensing authority for the quoted products (Hepatitis B) by the original manufacturer(s) to be submitted.
	- 1	BIS License with schedule for ISI Marked Products
		Quoted (wherever applicable) to be submitted. Average Annual Turnover Statement not less than 1 crores (One crore) of the bidder (Original
		Manufacturer / Importer) for Last 3 financial Years from Chartered Accountant with UDIN ANNEXURE-F to be submitted. Submitted turnover is without UDIN.

			ISO & CE/BIS/USFDA certificate for quoted Items as mentioned in bid Catalogue (wherever applicable) to be submitted. Submitted is not readable.
	M/s JK Diagnostics Rajkot (Manufacturer)		Average Annual Turnover Statement not less than 1 crores (One crore) of the bidder (Original Manufacturer / Importer) for Last 3 financial Years from Chartered Accountant with UDIN ANNEXURE-F to be submitted
			Copies of Audited Balance Sheet & Profit Loss Account for last three financial years from Chartered Accountant is not legible. Submit the readable copy of the same
			Valid Manufacturing License along with subsequent renewals, if any (K2EDTA) to be submitted
			Valid CGMP as per revised Schedule "M"/ WHO format to be submitted.
	±		Product permission by the licensing authority for the quoted products by the original manufacturer(s) to be submitted.
			Details of Technical personnel employed in the manufacturing and testing approved by the Licensing Authority to be submitted.
			BIS License with schedule for ISI Marked Products Quoted (wherever applicable) to be submitted.
			Latest Non Conviction Certificate issued by the Licensing authority of the respective state /UT (Issued not before 6 months) to be submitted.
			Non blacklisting / undertaking declaration(notarized) (Original manufacturer/ Direct Importer) on non-judicial Stamp paper of Rs 100/- (Annexure-F) to be submitted.
			Market Standing Certificate issued by Licensing authority of the respective States not Less than Three preceding years. (duly notarized) Annexure G to be submitted.
			ISO & CE/BIS/USFDA certificate for quoted Items as mentioned in bid Catalogue (wherever applicable) to be submitted. Submitted is not readable.

Note: - Firms are requested to submit all the documents well in time.

NO: JKMSCE/PS/44/KYON/7086-870

Copy for information to the:-

Managing Director JKMSCL

FA&CAO, JKMSCL

3. GM (Adm) JKMSCL

4. M/s Say Technologies for uploading of Notice on www.jkmsclbusiness.com

Office Record File.