Contract|अनुबंध

Contract No|अनुबंध क्रमांक: GEMC-511687718113373

Generated Date|अनुबंध तिथि : 09-Jan-2024

Organisation Details|संगठन विवरण

Type|प्ररूप :	Central Government
Ministry|मंत्रालय :	Ministry of Health and Family Welfare
Department|विभाग :	Department of Health and Family Welfare
Organisation Name|संगठन का नाम :	Central Health Service Ministry of Health and Family Welfare
Office Zone|कार्यालय क्षेत्र:	Safdarjung Hospital

Buyer Details|खरीदार विवरण

Designation|पद :	CMO
Contact No.|संपर्क नंबर :	011-26730507-
Email ID|ईमेल आईडी :	megha.arora@gov.in
GSTIN|जीएसटीआईएन :	-
Address|पता :	OFFICE OF THE MEDICAL SUPERINTENDENT SAFDARJUNG HOSPITAL NEW DELHI-110029, NEW DELHI, DELHI-110029, India

Financial Approval Detail|वित्तीय स्वीकृति विवरण

IFD Concurrence|आईएफडी सहमति:	Yes
Designation of Administrative Approval| प्रशासनिक अनुमोदन का पदनाम:	Medical Superintendent
Designation of Financial Approval| वित्तीय अनुमोदन का पदनाम :	Medical Superintendent

Paying Authority Details|भुगतान प्राधिकरण विवरण

Role:	DDO
Payment Mode| भुगतान का तरीका:	PFMS
Designation|पद :	ACCOUNTS OFFICER
Email ID|ईमेल आईडी :	acco@vmmc-sjh.nic.in
GSTIN|जीएसटीआईएन :	N
Address|पता:	OFFICE OF THE MEDICAL SUPERINTENDENT SAFDARJUNG HOSPITAL NEW DELHI-110029, NEW DELHI, DELHI-110029, India

Seller Details|विक्रेता विवरण

GeM Seller ID|जेम विक्रेता आईडी :	X8O5230008428595
Company Name|कंपनी का नाम :	DEXTRA MEDILINE
Contact No.|संपर्क नंबर :	07011113198
Email ID|ईमेल आईडी :	dextramedilineofficial@gmail.com
Address|पता :	4TH FLOOR, OFFICE NO. 405,KHASRA NO. 429,PROPERTY NO. 27,AGGARWAL CHAMBERS-4,,VEER SAVARKAR BLOCK, Shakarpur,Shahdara, East Delhi, DELHI-110092, -
MSME Registration number|एमएसएमई पंजीकरण संख्या :	-
GSTIN|जीएसटीआईएन:	07OGKPS8367C1ZL

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

Delivery Instructions|वितरण निर्देश : NA

Product Details|उत्पाद विवरण

#	Item Description|आइटम विवरण	Ordered Quantity| आइटम विवरण	Unit | इकाई	Unit Price (INR)| इकाई मूल्य (INR)	Tax Bifurcation (INR)| कर विभाजन (INR)	Price (Inclusive of all Duties and Taxes in INR)| मूल्य (INR में सभी शुल्क और कर सहित)
1	**Product Name|उत्पाद का नाम :** AQ 200 Excellus Fully Automatic Biochemistry Analyzer , Fully Automated Random Access System **Brand|ब्रांड :** AQ 200 Excellus **Brand Type|ब्रांड प्रकार :** Registered Brand **Catalogue Status|कैटलॉग की स्थिति:** OEM verified catalogue **Selling As|कैसे बेचा जा रहा है :** OEM verified Reseller **Category Name & Quadrant|श्रेणी का नाम और चतुर्थांश :** Fully Automatic Biochemistry Analyzer (V2) (Q2) **Model|मॉडल:** AQ 200 Excellus **HSN Code|एचएसएन कोड:** HSN not specified by seller	1	pieces	325,000	NA	325,000
Total Order Value |कुल ऑर्डर मूल्य (in INR)						**325,000**

Consignee Detail|परेषिती विवरण

S.No| क्र.सं.	Consignee|परेषिती	Item|वस्तु	Lot No.| लॉट नंबर	Quantity| मात्रा	Delivery Start After| दिनांक के बाद डिलीवरी शुरू करना है	Delivery To Be Completed By| वितरण पूरा कब तक करना है
	Designation|पद : CMO Email ID|ईमेल आईडी : megha.arora@gov.in	AQ 200 Excellus Fully				

1	Contact|संपर्क : 011-26730507- GSTIN|जीएसटीआईएन : - Address|पता : OFFICE OF THE MEDICAL SUPERINTENDENT SAFDARJUNG HOSPITAL NEW DELHI-110029, NEW DELHI, DELHI-110029, India	Automatic Biochemistry Analyzer , Fully Automated Random Access System	-	1	09-Jan-2024	24-Jan-2024

Product Specification for AQ 200 Excellus Fully Automatic Biochemistry Analyzer , Fully Automated Random Access System

Specification|विनिर्देश	Sub-Spec|उप-विनिर्देश	Value|मूल्य
GENERAL	Product Description	Fully Automatic Biochemistry Analyzer
	Purpose	Fully automatic biochemistry analyzer is used in clinical laboratories and healthcare settings to perform a wide range of biochemical tests on biological samples such as blood, serum, plasma, urine, and cerebrospinal fluid
PRODUCT INFORMATION	System Type	Fully Automated Random Access System
	Type of Biochemistry Analyzer	Bench Top Type
	Sample Type	Serum,Plasma,Whole Blood,Urine,CSF,Hemolyzed Blood,Any body fluid samples
	Assay Type	End point, Rate, Kinetic Turbidometric and Bi chromatic Assay
	Throughput tests / hour Photometric Tests with ISE module	200 or more
	Throughput tests/hour Photometric Tests without ISE Module	200 or more
	Calibration Facility	Factor,Linear (one, two and multi-point),Exponential,Logit-Log,Spline,Non-Linear
	Equipment must have self diagnostic tests with error message and online display	Yes
	Equipment must be programmable for all test menus and state of the art work station should be up-gradable by addition of ISE module	Yes
	Sample loading type	Continuous
	Equipment should have both internal and external probe cleaning/washing capacity	Yes
	Repeat facility to be available for calibrator and control	Yes
	Reagent refill message and monitoring facility	Yes
	Minimum number of test parameters on board at a time	≥ 30
	Minimum sample capacity at a time	35 to 49
	Minimum number of STAT samples at a time	≥ 10
	Should have pre post and auto dilution of samples and return capability for out of range samples	Yes
	Probe dispensers	Probe dispensers must have level detectors and must typically use between 2-25 µl of sample
	Spectoral range in nm	340 to 700 nm
	Wave length selection method	Interference Filter (With minimum 8 interference filter)
	Light source	Halogen lamp low cost with long life
	Life of lamp in hrs	≥ 1000
	Power consumption of lamp in VA	≤ 1000
	Type of sample cups	Reusable
	Number of sample cups provided with machine	1000
	QC program	QC Programme with L-J graphs, Print out of reports
	Reaction Cuvette	Reusable
	System should have on board cooling system	Refrigerated
	DI water plant for the washing of the cuvettes/ to avoid carry over and contamination	Yes
	Compatible on line UPS	Yes
	Back up time	1 hour
	Data management software	Equipment to be provided with compatible programmable windows based comprehensive data processing and management system graphical user interface software, LM Capability complete back up of data base for calibration control and patient sample results, capability for Bi-directional interfacing with LIS
	Patient result storage capacity	10000

	Onboard cuvette washing & drying facility	Yes
	Photometeric Checking of cuvettes before next reaction - Create Another field	Yes
	System should have facility for reading results on monitor and print out facility	Yes
	Reagents to be provided with machine	Creatinine,Urea,Uric Acid,Triglycerides,Glucose,Cholesterol,Total Protein,Calcium,Albumin,SGOT
	Quantity of Reagents provided (Number of Tests)	50 Tests or more
	Reports	Prinout of reports and full patient demographics to be available
	System to be provided with necessary pre requisites and start up kits normal and abnormal QC and calibrators	Yes
	Availability of RS 232 port/USB for data transfer	Yes
	Power supply	200-240V AC, 50Hz Single phase
	Reagents position	30 or more
	Bar code reader for reagents	Yes
	Bar code reader for samples	No
	Refrigerator provided with Analyzer	No
	Capacity of Refrigerator (Litres)	NA
	Type of Micro pipette provided	Not provided
	Variable Micro pipette provided	No
	Seller must provide original documentary proof of the date and place of manufacturing of provided equipment	Yes
	Service manual and operating manual to be provided	Yes
	Sytem shall be provided with a compatible desktop PC	Yes
	Speed of processor in GHZ	Greater than 3 GHZ
	RAM in MB	≥ 512 MB
	Hard disk drive capacity in GB	≥ 80 GB
	Monitor	TFT/LCD/LED/ Digital color monitor
	Monitor size (inches)	17 inches or more
	Compatible operating system provided in PC	Yes
	Printer provided	Yes
	Type of printer	Laser Printer
	The unit shall be capable of operating continuously in ambient temperature and relative humidity	Temp of 10° to 40° C and humidity of 15 to 90%
	Startup kit, Calibrators, consumables required in training the staff/Technician/Doctors should be provided at the time of installation &Training	Yes
	The reaction cuvettes should be replaced at free of cost when ever required during the warranty period	No
CERTIFICATION AND REPORTS	Compliance to Medical Device Rules (MDR) 2017 as amended till date	Yes
	Availability of valid Drug/medical device 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/Medical Device License Number	MFG/MD/2018/000022
	Manufacturing unit certification	ISO:13485 (Latest)
	Additional voluntary certification available	Q5 114470 0001 DATED- 14.03.2022
	Availability of Test Report for product as per Medical Device Rules (MDR) 2017 as amended till date	Yes
	Electrical Safety Standards	IEC/EN 60601-1 or equivalent BIS Standard
	Submission of all necessary certifications, licenses and test reports to the buyer at the time of bid submission and/or along with supplies as per buyer requirement	Yes
	OEM/Reseller shall ensure uninterrupted availability of all	

	spares for 10 years	Yes
	Availability of toll free facility for technical support maintened by OEM or authorized agencies	Yes
	User/Technical/Maintenance manuals to be provided in English in hard and soft copy	Yes
MISCELLANEOUS REQUIREMENTS	Details of equipments and procedures required for local calibration and routine maintenance to be provided and advanced maintenance task documentation also to be furnished	Yes
	List of important spares and accessories, with their part numbers to be provided to the buyer at the time of supplying the equipment	Yes
	Installation and Demonstration of equipment and training to be provided after completing supplies before acceptance	Yes
	The Principal Manufacturer must have direct Presence/approved service center In India	Yes
	Calibration certificates as per NABH requirement	Yes
WARRANTY	Warranty in years (including machine, DI Water Plant if provided , UPS, computer and printer)	3 year
ADDITIONAL REQUIREMENTS	Additional Requirements	NA

ePBG Detail | ईपीबीजी विवरण

NA

Terms and Conditions|नियम और शर्तें

1. Special terms and conditions- Version:1 effective from 20-09-2023

1.1

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 the submission of valid Drug License and self declaration of 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.**
6. **Comprehensive warranty: Comprehensive warranty shall include preventive maintenance including calibration as per technical/ service /operational manual of the manufacturer, service charges and spares. During the warranty period commencing from date of the successful completion of warranty period, Service personnel shall visit each consignee site as recommended in the manufacturer's technical/ service /operational manual, at least once in six months. warranty shall not be including the consumables. Further there will be 98% uptime warranty during warranty period on 24 (hrs) X 7 (days) X 365 (days) basis, with penalty, to extend warranty period by double the downtime period.**
7. **Service centres: Details of Service outlets in India to render services for equipment to be furnished to buyer/consignees with complete address, telephone numbers, e mails etc at time of making the supplies. It shall be the responsibility of seller to ensure that authorized service centres are available to cater to the areas where supplies are made within reasonable distance from where the service calls can be handled. Details of toll-free numbers for service call and online registration of service requests also to be provided buyer/consignee at the time of supplies.**
8. **Source of supply: It shall be responsibility of seller to provide Documents regarding source of equipments such as copy of Performa invoice or any other documents to establish that the products supplied are manufactured by OEM indicated and sourced from them.**
9. **Packing and Marking: Medical equipments being very delicate and sensitive packing for the goods should be strong and durable enough to withstand transit including transhipment (if any), rough handling, open storage etc. without any damage, deterioration etc. .The size, weights and volumes of the packing cases, remoteness of the final destination of the goods, availability or otherwise of transport and handling facilities at all points during transit up to final destination,. Quality of packing, the manner of marking within & outside the packages and provision of accompanying documentation shall take into consideration the type of medical equipments being supplied. The accessories shall be suitably labelled and packed. Each of the package shall be marked on three sides with indelible paint of proper quality: indicating contract number and date, brief description of goods including quantity, Packing list reference number, country of origin of goods and any other relevant details.**
10. **Spare Parts: Seller shall provide materials, information etc. pertaining to spare parts manufactured and supplied by the OEM. It shall be ensured that the required spares are available for purchase at least for 10 years from date of supplies. In case due to any reasons the production of the spare parts is discontinued sufficient advance notice should be given to the buyer/consignee before such discontinuation to provide adequate time to purchase the required spare parts etc. Further, OEM and their service centres/dealers shall carry sufficient inventories to assure ex-stock supply of consumables and spares for the equipments so that the same are available. OEM or reseller shall always accord most favoured client status to the buyer/consignee and shall give the most competitive price for spares and consumables of its machines/equipments supplied.**
11. **Installation, Training, Manuals: Seller shall be responsible to carry out Installation & commissioning, Supervision and Demonstration of the goods. They shall provide required jigs and tools for assembly, minor civil works for the completion of the installation and Training of Consignee's representatives for operating and**

maintaining the equipment and supplying required number of operation & maintenance manual for the goods. In case the category parameters are specifying any requirements regarding the installations, training and manuals the same shall also be applicable.

12. Electrical safety checking: Sellers are required to make sure that they furnish the list of equipments for carrying out routine and preventive maintenance to buyer/consignee .They should make sure to periodically check the electrical safety aspects as per BIS Safety Standards or equivalent .In case they do not have required equipment for such testing should ensure that the equipments checked for electrical safety compliance through labs with facilities for such checking during every preventive maintenance call.
13. Software: All software updates should be provided free of cost during warranty period.

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