

अनुबंध | Contract



अनुबंध क्रमांक | Contract No: GEMC-511687766980581

अनुबंध तिथि | Generated Date : 26-Apr-2024

संगठन विवरण Organisation Details		खरीदार विवरण Buyer Details				
प्ररूप Type :	Central Autonomous	पद Designation :	Assistant Administrative Officer			
मंत्रालय Ministry :	Ministry of Health and Family Welfare	संपर्क नंबर Contact No. :	0535-2979749-1103			
विभाग Department :	Department of Health and Family Welfare	ईमेल आईडी Email ID :	buycon38.aiimsa.up@gembuyer.in			
संगठन का नाम Organisation Name :	All India Institute of Medical Sciences (AIIMS)	जीएसटीआईएन GSTIN :	09AAAJA2765Q1ZS			
कार्यालय क्षेत्र Office Zone :	AIIMS Raebareli Uttar Pradesh	पता Address :	All India Institute Of Medical Sciences, Munshiganj, Dalmau Road, Raebareli, RAEBARELI, UTTAR PRADESH-229405, India			
वित्तीय स्वीकृति विवरण Financial Approval Detail		भुगतान प्राधिकरण विवरण Paying Authority Details				
आईएफडी सहमति IFD Concurrence :	Yes	Role:	PAO			
प्रशासनिक अनुमोदन का पदनाम Designation of Administrative Approval:	DDA, AIIMS, Raebareli	भुगतान का तरीका Payment Mode:	Offline			
वित्तीय अनुमोदन का पदनाम Designation of Financial Approval :	Financial Advisor	पद Designation :	ACCOUNTS OFFICER			
		ईमेल आईडी Email ID :	ddo1.aiims.up@gembuyer.in			
		जीएसटीआईएन GSTIN :	-			
		पता Address:	All India Institute Of Medical Sciences, Munshiganj, Dalmau Road, Raebareli, Raebareli, UTTAR PRADESH-229405, India			
विक्रेता विवरण Seller Details						
जेम विक्रेता आईडी GeM Seller ID :	5LIE230009090775					
कंपनी का नाम Company Name :	OSHNAM BIOTECH					
संपर्क नंबर Contact No. :	09179055137					
ईमेल आईडी Email ID :	oshnabiz@gmail.com					
पता Address :	LGF , 1,1Garhi Mutwali, Adjacent Bank of India, Dalmau Road, Munshiganj, Raebareli, UTTAR PRADESH-229405, -					
एमएसएमई पंजीकरण संख्या MSME Registration number :	-					
जीएसटीआईएन GSTIN:	09AAIFO0381Q1ZN (R)					
*जिसके नाम के पक्ष में GST/TAX इनवॉइस पेश किया जाएगा GST / Tax invoice to be raised in the name of - Buyer						
वितरण निर्देश Delivery Instructions : NA						
उत्पाद विवरण Product Details						
#	आइटम विवरण Item Description	आइटम विवरण Ordered Quantity	इकाई Unit	इकाई मूल्य (INR) Unit Price (INR)	कर विभाजन (INR) Tax Bifurcation (INR)	मूल्य (INR में सभी शुल्क और कर सहित) Price (Inclusive of all Duties and Taxes in INR)
1	उत्पाद का नाम Product Name : bpl cardiart ECG Machine 12 Channel ब्रांड Brand : bpl cardiart ब्रांड प्रकार Brand Type : Registered Brand कैटलॉग की स्थिति Catalogue Status: OEM verified catalogue कैसे बेचा जा रहा है Selling As : OEM verified Reseller श्रेणी का नाम और चतुर्थांश Category Name & Quadrant : Electrocardiography (ECG) Machine (V2) (Q2) मॉडल Model: CARDIART 9108 एचएसएन कोड HSN Code: HSN not specified by seller	2	pieces	155,000	NA	310,000
कुल ऑर्डर मूल्य Total Order Value (in INR)						310,000
परोक्षी विवरण Consignee Detail						
क्र.सं. S.No	परोक्षी Consignee	वस्तु Item	लॉट नंबर Lot No.	मात्रा Quantity	दिनांक के बाद डिलीवरी शुरू करना है Delivery Start After	वितरण पूरा कब तक करना है Delivery To Be Completed By
1	पद Designation : Assistant Administrative Officer ईमेल आईडी Email ID : buycon38.aiimsa.up@gembuyer.in संपर्क Contact : 0535-2979749-1103 जीएसटीआईएन GSTIN : 09AAAJA2765Q1ZS	bpl cardiart ECG	-	2	26-Apr-2024	11-May-2024

पता | Address : All India Institute Of Medical Sciences, Munshiganj,
Dalmau Road, Raebareli,
RAEBARELI, UTTAR PRADESH-229405, India

Machine 12 Channel

Product Specification for bpl cardiart ECG Machine 12 Channel

विनिर्देश Specification	उप-विनिर्देश Sub-Spec	मूल्य Value
GENERAL FEATURES	Product Description	ECG Machine
	Purpose	An ECG is a noninvasive routine examination of the electrical activity of the heart that is used to reflect underlying heart conditions
PRODUCT INFORMATION	Operating modes of ECG Machine	Automatic, Manual and Rhythm
	ECG machine should have ECG lead annotation facility	Yes
	Leads which is in ECG machine should be able to acquire simultaneously and interpret them	12
	Number of channels	12 Channel
	ECG machine should acquire lead ECG for both adult and pediatric patients	Yes
	The ECG machine should have facility to show lead fail indication	Yes
	The ECG machine should have facility to show lead reversal indication	Yes
	The ECG machine should have facility to show the impedance to quality check of connection	Yes
	Acquisition time for ECG Machine in sec	10 sec
	Digital sampling rate for Pacemaker spike detection	8000 s/sec/channel
	Recording of digital sampling for pacemaker	1000 s/sec/channel
	ECG machine should have real time colour backlit display of ECG waveforms with signal qualify indication for each lead	Yes
	ECG machine should have frequency filters	Artifact, AC and low and high pass frequency filters
	Number of ECGs which can be store in ECG Machine	151 to 200
	ECG machine should have full screen preview of ECG report for quality assessment checks prior to print	Yes
	Type of inbuilt screen	LCD
	Size of screen in inches	5
	Display resolution of ECG machine in pixels	640 x 480
	ECG machine should have interpretation facility of the amplitudes, duration and morphologies of ECG waveforms and associated rhythm for adult and pediatric patient	Yes
	ECG machine should have alphanumeric keyboard for patient data entry	Hard keys
Availability of latest interpretation software	No	
Wireless acquisition module with RF technology	No	
System should have the dedicated software to download the ECG form machine in PDF format	Yes	
PRINTER	Printer Type	Thermal Printer
	Recorder Paper Size	A4 Size
	Recorder Speed	50 mm/sec
	Resolution of digital array printer	200 dpi x 500 dpi
	Number of Thermal paper	500
	ECG machine report format	Report formats of 3x4, 6x2, Rhythm for up to selected leads, 12 lead extended measurement, 1 minute of continuous waveform data for 1 selected lead
BATTERY	Provision of Battery	Yes
	Battery Type	Built in Rechargeable Battery
	Battery capacity of continuous rhythm recording on single charge (minutes)	60
	Battery capacity	50 ECG or 1 hour of continous rhythm recording on single charge
	Connectivity to ECG Machine	LAN
	Storage on external portable memories	USB support
	The individual patient lead should be change without replacing the whole patient	Yes

	cable assembly	
ENVIRONMENTAL PARAMETERS	Operating temperature and humidity	Temperature of 10 to 40 degree Celsius and Relative Humidity of 15 to 90%
POWER SUPPLY	Power input	220-240 V AC, 50 Hz fitted with Indian plug
ACCESSORIES	ECG Machine 12 leads with interpretation	1
	Patient Cable	2
	Chest Electrodes Adult (set of Six)	1
	Chest Electrodes Paediatric (set of Six)	Not provided
	Limb Electrodes	2 for adults and 2 for paediatric
	Power cable for charging	Yes
	Supplied with Clip electrode	Yes
	Compatible trolley provided	No
CERTIFICATION & REPORTS	Compliance to Medical Device Rules (MDR) 2017 as amended till date	Yes
	Availability of valid drug 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 License Number	NA
	Manufacturing unit certification	ISO:13485 (Latest)
	Additional voluntary certification available	NA
	Availability of Test Report for each supplied batch/product as per Medical Device Rules (MDR) 2017 as amended till date	Yes
	Compliance to Safety Standards	IS 13450:Part 2:Sec 25 / IEC 60601-2-25
	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
WARRANTY	Warranty (Option of comprehensive warranty is available through bidding only, which if opted will supersede normal warranty in the catalogue)	2 year
ADDITIONAL REQUIREMENTS	Additional Requirements	NA
Miscellaneous Parameters	User/Technical/Maintenance manuals to be supplied in English in hard and soft copy	Yes
	The Principal Manufacturer must have direct Presence/approved service center In India	Yes
	Installation and demonstration of equipment and training to be provided after completing supplies before acceptance	Yes

ईपीबीजी विवरण | ePBG Detail

NA

नियम और शर्तें | Terms and Conditions

1. Special terms and conditions- Version:1 effective from 23-08-2023

- 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.
 - The sellers are registered on GeM based on self-declaration of valid Drug License, 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.
 - 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.
 - 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.
 - 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.
 - 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 transshipment (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

नोट: यह सिस्टम जनरेटेड फाइल है। कोई हस्ताक्षर की आवश्यकता नहीं है। इस दस्तावेज़ का प्रिंट आउट भुगतान/लेनदेन उद्देश्य के लिए मान्य नहीं है।

Note: This is system generated file. No signature is required. Print out of this document is not valid for payment/ transaction purpose.