

Contract | अनुबंध



Contract No | अनुबंध क्रमांक: GEMC-511687715736546

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Organisation Details संगठन विवरण Type प्ररूप : State Government Ministry मंत्रालय : - Department विभाग : Medical Health and Family Welfare Department Uttar Pradesh Organisation Name संगठन का नाम : Combined Admission System (CAS), Department of Medical Health & Family Welfare, Uttar Pradesh Office Zone कार्यालय क्षेत्र : Cmo Mathura		Buyer Details खरीदार विवरण Designation पद : NODAL OFFICERS Contact No. संपर्क नंबर : 0565-2471914- Email ID ईमेल आईडी : buycon13.cascdup.up@gembuyer.in GSTIN जीएसटीआईएन : - Address पता : CMO Office Mathura, MATHURA, UTTAR PRADESH-281001, India				
Financial Approval Detail वित्तीय स्वीकृति विवरण IFD Concurrence आईएफडी सहमति : No Designation of Administrative Approval प्रशासनिक अनुमोदन का पदनाम : cmo Designation of Financial Approval वित्तीय अनुमोदन का पदनाम : cmo		Paying Authority Details भुगतान प्राधिकरण विवरण Role : PAO Payment Mode भुगतान का तरीका : Offline Designation पद : lipik Email ID ईमेल आईडी : pao2.hd.mathura@gembuyer.in GSTIN जीएसटीआईएन : - Address पता : CMO Office Mathura, MATHURA, UTTAR PRADESH-281001, India				
Seller Details विक्रेता विवरण GeM Seller ID जेम विक्रेता आईडी : 3788180000473850 Company Name कंपनी का नाम : M/S SWAASTI ENTERPRISES Contact No. संपर्क नंबर : 09140711660 Email ID ईमेल आईडी : swaastienterprises17@gmail.com Address पता : Villa No-B, 2,Shalimar Garden Bay,IIM Road,Luckno, Lucknow, UTTAR PRADESH-226020, - MSME Registration number एमएसएमई पंजीकरण संख्या : - GSTIN जीएसटीआईएन : 09AVCPT7930K1ZZ						
*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 : W WARSIMPEX MEDICAL SYSTEMS INDIA quartz infrared Clinical infant warmers Brand ब्रांड : W WARSIMPEX MEDICAL SYSTEMS INDIA Brand Type ब्रांड प्रकार : Registered Brand Catalogue Status कैटलॉग की स्थिति : Catalogue not verified by OEM Selling As कैसे बेचा जा रहा है : Reseller not verified by OEM Category Name & Quadrant श्रेणी का नाम और चतुर्थांश : Clinical incubators or infant warmers (Q3) Model मॉडल : WARSIMPEX -R-W HSN Code एचएसएन कोड : HSN not specified by seller	6	pieces	60,000	NA	360,000
Total Order Value कुल ऑर्डर मूल्य (in INR)						360,000
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
1	Designation पद : acmocmsdstore Email ID ईमेल आईडी : dycmo.au-up@gov.in Contact संपर्क : - GSTIN जीएसटीआईएन : - Address पता : CMO Office Mathura,	W WARSIMPEX MEDICAL SYSTEMS INDIA quartz infrared Clinical infant warmers	-	6	29-Dec-2023	13-Jan-2024

Product Specification for W WARSIMPEX MEDICAL SYSTEMS INDIA quartz infrared Clinical infant warmers

Specification विनिर्देश	Sub-Spec उप-विनिर्देश	Value मूल्य
Technical Characteristics (specific to this type of device)	facility to display skin set, skin observed temperature in degree C and heat power separately	No
	have user friendly touch panel control	Yes
	Type of heater:	quartz infrared
	Audiovisual alarm facility for:	Overheating beyond set temperature range, Spurt in Heater output i.e. more than 60% heater output for 10 minutes system should raise alarm and cutoff heater, Power failure, Heater failure, Patient temperature less/Greater than required/set temperature, Probe failure
	Warmer head should be rotatable in different direction, so as to allow taking X-ray	Yes
	Observation light Luminance and Colour temperature	> = 1000 Lux , Colour temperature range from 3700K to 5100K
	Battery back up facility for Power failure indication during power fail	Yes
	desired temperature range	25 to 40 degree C
	settable temperature 32 to 38 deg C	32 to 38 deg C
	temperature resolution should be:	<= 0.1 degree C
	Temperature accuracy should be	<= 0.2 degree C
	facility to lock the keyboard to avoid unwanted user modification of the set parameters	Yes
	should have separate Bassinet trolley, bed should be tilt-able and have suitable provision for x-ray cassette holder, Mattress foam density should be minimum 25 kg/cm ³ , transparent collapsible side walls easily detachable for cleaning	Yes
	Should have a Feather Touch operation with large digital display and comprehensive alarms, Control Panel should be liquid proof and allow easy and hygienic disinfection	Yes
	Manual Mode can adjust Heater Output 10 -100 %, with 10% increment, an auditory and visual alarm shall be given at least every 15 min	Yes
	Under manual mode, heater cut off / switch off , if the maximum irradiance at any point of the mattress area exceeds a total irradiance level of 10 mW/cm ² (between 10 to 30 minutes)	Yes
	Bed height from Floor	81 centimeter
	Bed Distance from the heat source	90 centimeter
	should have lockable castor wheels	Yes
	Green indicator light shall be indicate that warmer is ready for normal use	Yes
	Markings on the bassinet and X-Ray cassette holder to enable proper positioning of the baby while doing the X-Ray	Yes
	Number of tubing ports (edges covered by silicon rings) on the side walls (For cable management)	2
	Height of the side walls over the mattress	110 millimeter
	Provision of X-Ray cassette tray of at least 750 mm X 350 mm size, suitable to adopt <= 20mm thick X-Ray cassette	Yes

	bay bed should be crevice free for ease of cleaning, infection control	Yes
	mattress used should be of biocompatible material	Yes
	Skin temperature probe should be small in size [Diameter <= 10 mm & Height <= 4 mm] with biocompatible Baby contact material	Yes
	Mobility, portability	On castors (2 of the castors should have brakes) & Castor size >= 4 inch.
Definition and Clinical Purpose	Definition	Mains electricity (AC-powered) mobile device that contains an infrared (IR) heating element(s) designed to emit controlled, evenly distributed overhead heat to the body of a newborn/infant patient requiring supplemental heat. This device is equipped with wheels so that it can easily be moved to different areas of a room, ward, or department.
	Clinical purpose & Overview of functional requirements	It is an electrically powered device with a radiant heating source intended to maintain the thermal balance of an infant by direct radiation of energy in the infrared region of the electromagnetic spectrum. It is a microprocessor controlled unit with heater placed on the over head panel. This work on both servo and manual mode options to maintain the baby temperature at the set value. There are two modes of operation manual and baby (servo) mode. It has Digital displays reading the set and baby observed temperatures separately.
Settings	Settings with facility to clearly display the selected mode, Option of Selecting either Manual mode and Baby (Servo) mode settings, Provision for Set temperature range (in servo mode) from 32 to 38 deg C	Yes
	User's interface	Manual and Servo controlled temperature regulation
	Software and/or standard of communication (where ever required)	LED Display and inbuilt software; Interruption and restoration of the power supply does not change the preset values
	Others:	1. Device shall not overbalance when placed in any transport position of normal use on a 10° inclined plane from the horizontal plane. 2. Transformers of device shall be protected against overheating in the event of short circuit or overload of any output winding. 3. Patient leakage current should be less than 100 µA in normal condition 4. Temperature on the baby mattress should not exceed 43 deg C when the warmer is operating under steady temperature condition 5. Temperature of HEATER GUARDS should not exceed 85 °C in normal use. 6. The Temperature differences on the mattress shall not exceed 2 °C.
PHYSICAL CHARACTERISTICS	Overall Height of equipment	1500 millimeter
	Overall Width of equipment	900 millimeter
	Overall Length of equipment	700 millimeter
	Overall weight of equipment	150 kilogram
	Heat source Configuration	>= 60 degree angle adjustment must be possible in the heat source, it should provide shielding to the infant in case of breakage of tubes/bulbs and all surfaces to be made of corrosion resistant material
	Noise	Auditory alarm shall have a sound level of at least 65 dBA at a distance of 3 m from the front of the infant radiant warmer, and the sound level of the alarm shall not exceed 80 dBA on the mattress
	Heat dissipation	Should maintain desired set temperature and uniform heat distribution in baby cradle
ENERGY SOURCE	Power Requirements	220 to 240V, 50 Hz AC, ± 10% of input
	Battery operated	Power failure indication during power fail
	Protection	OVP, earth leakage protection
	Power consumption	701 Watt
	Compatible to alternate energy supply source; Solar Heating in addition to AC power source):	No
	Thermal reflector to fix the skin probe	>= 50 units to be supplied with each equipment unit
ENVIRONMENTAL AND DEPARTMENTAL CONSIDERATIONS	Atmosphere / Ambiance	Operating condition: -Capable of operating continuously in ambient temperature of 0 to 50 deg C and relative humidity of 15 to 90% in ideal circumstances. - an ambient air velocity is less than 0.3 m/s.
	User's care, Cleaning, Disinfection & Sterility issues	Complete unit to be easily washable and sterilizable using both alcohol and chlorine agents.
STANDARDS AND SAFETY	Certificates,	CE European
	Performance and safety standards (specific to the device type); Local and/or international	IEC-60601-1-2:2007 Medical electrical equipment Part 1-2: General requirements for basic safety and essential performance - Collateral standard: Electromagnetic compatibility -Requirements and tests (Or Equivalent BIS). Shall meet IEC 60601 -2- 21: 2009 Medical Electrical Equipment - Part 2-21: Particular Requirement for the basic safety and essential performance of infant radiant warmers . should meet IEC 60601-1:2005(or latest) standard requirements. Baby contact material should be biocompatible. Manufacturer should be ISO 13485 certified
	Availability of Type Test report from any Govt/ NABL accredited /ILAC approved laboratory (To be produced by Seller on	Yes

	Buyer's demand if indicated available)	
TRAINING AND INSTALLATION	Pre-installation requirements for the equipment	Availability of 5 amp/15 Amp. Electrical socket (2 nos) for each warmer at Buyer's facility.
	Documentation required to be provided by Seller:	User & Operating manual, Certificate of Calibration, Factory inspection report, user training manual
WARRANTY	Warranty (Option of comprehensive warranty is available through bidding only, which if opted will supersede normal warranty in the catalogue)	3 year

Note | टिप्पणी: Seller has given an undertaking that it has made arrangements for getting the stores from an authorized distributor / dealer / channel partner of the OEM of the offered product. At the time of delivery of goods, Seller will provide necessary chain documents (in the form of GST Invoice) to prove that the supplied goods are genuine and are being sourced from an authorized distributor / dealer / channel partner of the OEM. In case of any complaint about genuineness of the supplied products, Seller shall be responsible for providing genuine replacement supplies.

ePBG Detail | ईपीबीजी विवरण

NA

Terms and Conditions | नियम और शर्तें

1. Special terms and conditions- Version:2 effective from 11-11-2022

1.1

- All Provisions of Drugs and Cosmetics Act, 1940 and Rules made there under as amended till date will always be applicable. This will include all notifications issued by Central Drugs Standard Control Organisation (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.
- The price offered by the seller shall not, in any case exceed the DPCO controlled price, if any, fixed by the Central/State Government, the Maximum Retail Price (MRP) and the selling price. The seller must reduce the prices if there is any reduction in DPCO ceiling price, if any.
- 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.
- 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.
- 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.
- 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.
- 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.
- 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.
- 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.
- Software: All software updates should be provided free of cost during warranty period.
- 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 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.

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

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

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।