	अनुबंध Сс	ontract							
<u>-</u>	GeM 7		अनुबंध	ध क्रमांक (	Contract No: G	EMC-5	116877884776	15	
Efficient - Transpa	Azadi Ka Amrit Mahotsav		अनुबंध तिथि  Generated Date : 17-Jun-2025						
संगठन विवेर	ण Organisation Details	ख	वरीदार विवरण	Buyer I	Details				
प्ररूप Type :	तरूप Type : Central Autonomous नंत्रालय Ministry : Ministry of Health and Family Welfare वभाग Department : Department of Health and Family Welfare		द Designation पर्क नंबर Contac	:	Stores and Procurment Officer				
संगठन का			ईमेल आईडी Email जीएसटीआईएन GS	STIN :	harendra.dey@nic.in N P.O. NEIGRIHMS, Mawdiangdiang, Shillong, KHASI HILLS EAST, MEGHALAYA-793018, India			_	
:			ता Address :						
<u> </u>	~	-		0					
•	ति विवरण Financial Approval Detail				Paying Auth	ority	Details		
	ाईएफडी सहमति]IFD Concurrence : Yes शासनिक अनुमोदन का पदनाम		Role: भुगतान का तरीका		ΡΑΟ				
Designation	ination of Administrative Director Concurrence C8491of2025and2026		Payment Mode:		Offline				
Approval: वित्तीय अनुमोदन का पदनाम  DDA & DFA c849 of13.6.2025 GIA- Asset within 1 Designation of Financial Approval : 5 days			द Designation मेल आईडी Email ोएसटीआईएन GS	ID :	Thwet Star Syngkon thwet.syngkon@neigrihms.gov.in				
Designation of Financial Approval : 5 days			पता। Address:		- P.O. NEIGRIHMS, Mawdiangdiang, Shillong, KHASI HILLS EAST, MEGHALAYA-793018, India				
<sup>जीएसटीआईएन</sup> *जिसके ना वितरण निवै under G	करण संख्या MSME Registration number : -  GSTIN: 27AAACI4227Q1Z8 (R), (B)  म के पक्ष में GST/TAX इनवॉइस पेश किया जाएगा GST / Ta ईश   Delivery Instructions : Processing of Dual C IA-Asset-25-26 for department of Cardiology ग Product Details आइटम क्विरण Item Description					h thr		ranty मूल्य (INR में सभी शुल्क और क सहित)  Price	
उत्पाद का नाग	म   <b>Product Name :</b> medtronic Dual Chamber Temporary Pacemaker		Quantity		(INR)		(INK)	(Inclusive of all Duties ar Taxes in IN	
ब्रांड प्रकार B कैटलॉग की वि कैसे बेचा जा श्रेणी का नाम (Q2) मॉडल Mod	l : medtronic Brand Type : Registered Brand स्थिति Catalogue Status: OEM verified catalogue रहा है Selling As : OEM और चतुर्यांश Category Name & Quadrant : Dual Chamber Temporary Pac lel: EPG-5392 Dual Chamber Temporary Pacemaker ls HSN Code: HSN not specified by seller	cemaker	2	pieces	250,000	NA		500,000	
 ज्ल ऑर्डर मूल्य	Total Order Value (in INR)					ļ		500,000	
पराषता विवर	ण Consignee Detail						[	विकाण प्राय क	
	परेषिती  Consignee	वस्तु	Item	लॉट नंब Lot N	' Hiari Ouz	antity	दिनांक के बाद डिलीवरी शुरू करना है  Delivery Start	वितेरण पूरा क तक करना है  Delivery To Be	
क्र.सं. S.No							After	Completed By	

विनिर्देश Specification	। उप-विनिर्देश Sub-Spec	मूल्य Value						
	Product Description	Dual Chamber Temporary Pacemaker						
GENERAL	Purpose	A dual chamber temporary pacemaker is a small, portable device that is used to regulate the heart rhythm.						
	Modes	DDD,DOO,DDI,AAI,AOO,VVI,VOO						
	Pacing Rates	30 to 200 ppm						
	Upper Rate	80 to 230 ppm						
	Rapid Atrial Pacing Rates	80 to 800 ppm						
PRODUCT INFORMATION	Defibrillator Protection	Yes						
	Continuous monitoring of the battery voltage	Yes						
	Backup pacing during battery change	Yes						
	Battery Type			Replacable with non rechargable batteries				
	Lock protection for the parameters through physical or software options				Yes			
	Compliance to Medical Device Rules (MDR) 2017 as amended till	Yes						
	Availability of valid 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 Medical Device License Number	2697-12-2023						
CERTIFICATIONS	Certification for manufacturing unit	ISO:13485 (Latest)						
	Availability of Test Report for each supplied batch/product as pe 2017 as amended till date	Yes						
	Submission of all necessary certifications, licenses and test repo bid submission or along with supplies as per buyer requirement	Yes						
	Conformity to standard	IEC 60601-2-31 or Equivalent BIS						
WARRANTY	Warranty (Years)			3 year				
ADDITIONAL REQUIREMENTS	Additional Requirements			NA				
ईपीबीजी विवरण   eP								

## नियम और शर्तें|Terms and Conditions

1. Special terms and conditions- Version:1 effective from 15-03-2024

- 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.
  - The sellers are registered on GeM based on the self declaration of valid Medical Device 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.
  - 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
- 2.3 All GeM Sellers / Service Providers are mandated to ensure compliance with all the applicable laws / acts / rules including but not limited to all Labour Laws such as The Minimum Wages Act, 1948, The Payment of Wages Act, 1936, The Payment of Bonus Act, 1965, The Equal Remuneration Act, 1976, The Payment of Gratuity Act, 1972 etc. Any non-compliance will be treated as breach of contract and Buyer may take suitable actions as per GeM Contract.

नोट: यह सिस्टम जनरेटेड फाइल है। कोई हस्ताक्षर की आवश्यकता नहीं है। इस दस्तावेज़ का प्रिटे 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.