



1	ईमेल आईडी   Email ID : dr.brahmdeo.kumar@esic.nic.in संपर्क   Contact : 091-93863685-63 जीएसटीआईएन   GSTIN : 10PTNE01028G1D7 पता   Address : ESIC HOSPITAL BIHTA NH-30, SIKANDERPUR PATNA ARA ROAD, BIHTA PATNA-801103 LANDMARK- ADJACENT TO NDRF CAMPUS BIHTA, PATNA, BIHAR-801103, India	ORIKAM Dental X-Ray Machine	-	1	07-Jun-2024	22-Jun-2024
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### Product Specification for ORIKAM Dental X-Ray Machine

विनिर्देश   Specification	उप-विनिर्देश   Sub-Spec	मूल्य   Value
GENERAL	Product Name	Dental X-Ray Machine
	Purpose	Dental X-ray machines are used to capture images of the teeth, jawbones, and surrounding oral structures
PRODUCT INFORMATION	Tube voltage (KV)	65 KV
	Availability of selection option for Bisecting angle technique & Parallel Technique	Yes
	Tube current (mA)	2 to 8 mA
	Focal spot (mm)	0.4 to 0.8 mm
	Should have Total filtration and Inherent Filtration	Yes
	Total filtration in mm Al	≥ 2 mm Al
	Inherent filtration	≥ 0.5 mm Al
	Minimum Exposure Time	0.02 to 2 seconds
	Power supply	220-240 V, 50 Hz single phase
	Based on DC Current	Yes
	RVG / X Ray Film compatible	Yes
	Machine should have a stable base with sturdy locking wheel mechanism	No
	Wall Mountable	No
	Digital control equipped with an easy ready display indicating with precision	Yes
	Stool for the patient provided	No
	Soft positioning arms for accurate tube positions	Yes
	Light and flexible movements	Yes
Internally lead coated head tube and cone to avoid scattered radiations	Yes	
Exclusive angular indicating system for head positioning in various radiography techniques	Yes	
High efficiency and greater sharpness of the radiography	Yes	
Additional Parameters	Should have high voltage generator with high efficiency in the emission of the X rays	Yes
	Should have free swivel head, which allows easy positioning of head	Yes
	Number of lead aprons provided	2 (0.5 mm lead equivalent)
	To be supplied with Thyroid shield	Yes
CERTIFICATION AND 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/Medical Device License Number	MFG/MD/2023/000772
	AERB Type Approved Model	Yes
	AERB approval certificate number of Particular Model	HR-94853-MF-XRE-001
	Additional voluntary certification available	HR-94853-MF-XRE-001
	Conformity to Indian Standard IS 7620- Part 1, Part 2, Part 3	Yes
	Manufacturing unit certification	ISO:13485 (Latest)
	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
WARRANTY	Warranty (Option of comprehensive warranty is available through bidding only, which if opted will supersede normal warranty in the catalogue)	4 year
	User/Technical/Maintenance manuals to be supplied in English in hard and soft copy	Yes

MISCELLANEOUS REQUIREMENTS	Installation and Demonstration of equipment and training to be provided after completing supplies before acceptance	Yes
	Principal Manufacturer must have direct Presence/approved service center In India	Yes
	Availability of toll free facility for technical support maintained by OEM or authorized agencies	Yes
	OEM/Reseller shall ensure uninterrupted availability of all spares for 10 years	Yes
ADDITIONAL REQUIREMENTS	Additional Requirement	NA

उत्पाद का बीआईएस लाइसेंस | BIS license of the product: 9512474924

### ईपीबीजी विवरण | ePBG Detail

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### नियम और शर्तें | Terms and Conditions

#### 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 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।

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