

# REPUBLIC OF THE PHILIPPINES Department of Health

### NATIONAL CENTER FOR MENTAL HEALTH

Nueve de Febrero Street, Mandaluyong City, Philippines



#### **BIDS AND AWARDS COMMITTEE**

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## SECTION VI SCHEDULE OF REQUIREMENTS

#### ITB No. 044-2023

The delivery schedule expressed as weeks/months stipulates he reafter a delivery date which is the date of delivery to the project site.

ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
US23-01	FLOOR MOUNTED DIGITAL X-RAY MACHINE High Voltage Generator  1. Maximum Output Power: 30 kW or better  2. Maximum Exposure Voltage: 120 kV or better  3. Maximum mA: 400 mA or better  4. Power requirements: 220V Three-Phase with Isolation transformer if necessary  X-ray Tube Assembly	1	11,000,000.00	90 calendar days
	<ul> <li>5. Dual Focal Spot</li> <li>6. Rotating Anode</li> <li>Floor Mounted Tube Stand with Patient Table</li> <li>7. At least four-way table</li> </ul>			
	Digital Detector  8. Detector type: Amorphous Silicon TFT / Cesium Iodide scintillator (CsI)  9. Auto detect location of detector  10. Detector Size: 14 x 17 inches  11. Connection: Wireless with 2 extra battery			
	12. Image Acquisition time: at least 5 sec. 13. Dust and water resistant (at least IP55)  Vertical Bucky Stand 14. Grid compatible to the detector			

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	15. Auto sync/tracking feature between tube and vertical bucky			
	Patient Table 16. Auto tracking: Longitudinal tracking of detector with longitudinal travel and rotation of tube.			
	Clinical Applications 17. With DAP meter for patient radiation exposure			
	Data transfer  18. DICOM 3.0 compliant  19. Support digital data transfer via a DICOM network for both printing & storage.  20. Shall be capable of sending DICOM images to our PACS/RIS server  21. The system shall have a Open Modality Worklist Function to be connected to our existing PAC/RIS.			
	Standard Accessories  22. Three (3) Full body Lead apron  23. Two (2) thyroid shields  24. One (1) upright gonad shield  25. One (1) upright gonad shield  26. Two (2) movable lead barriers  27. QA Phantom  a. Collimation and beam alignment test  b. Spatial Resolution test  c. Contrast Resolution test  d. Uniformity test			
	Inclusions  28. Four (4) workstations with at least;  a) Processor: at least Intel Core i7 (11 <sup>th</sup> gen)  b) Memory: 8 GB RAM c) Hard Disk: 1 TB HDD d) Operating System: Windows 10 OS and license Microsoft Office e) Video cards compatible to the existing PACS/RIS.			

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	29. Each workstation will be supplied with keyboard, mouse, Dual monitor (24" and 17") LED HD monitors, UPS and AVR 30.2 pieces 4-Seater Gang Chair 31.3 units of air purifier with H13 filter for a 40 sqm room 32.10 extra H13 filters 33. One (1) UV sterilizer for a 40 sqm room 34. One (1) All-in-one Heavy duty printer with 10 extra inks 35. One (1) 3 tonner tower type Aircon (inverter type) 36. Two (2) window type inverter Aircon (1.5 HP) 37. Lead glass (14x14) 38. Lead Shielding to the examination room if needed 39. AVR for Digital Xray machine 40. One (1) Mobile X-ray table with bucky 41. One (1) Mobile Cassette Holder (14 x 17) 42. With civil and electrical works needed for the X-ray examination room and			
	<ul> <li>Warranty</li> <li>43. With three (3) years warranty for parts and services with free quarterly preventive maintenance after end user acceptance. Warranty shall commence upon the acceptance of the end user. The winning bidder shall issue a certificate reflecting the duration of warranty period.</li> <li>44. Bidder's certificate guaranteeing the availability on the supply of spare parts ten (10) years from end of production. A certification must be issued by the manufacturer for the bidder.</li> <li>45. With quarterly preventive maintenance and calibration</li> <li>46. Certification in case that the equipment will be for repair or maintenance within the warranty period, it should be done on-site.</li> </ul>			

Standard Requirements  47. The supplier shall secure a certificate from the manufacturer stating that the equipment is brand new, unused and not a discontinued model or was listed in the market recall. The certificate reflect the brand, model, serial number of the machine, and the date and place it was manufactured.  48. System must not be a retrofit solution Detector and x-ray machine are of the same brand.  49. The bidder must provide the current and valid certificate of Manufacturer's compliance with ISO 13485.  50. Supplier/ Bidder must be ISO certified  51. Delivery of equipment and all accessories within 90 calendar days	ITEM NO.	DESCRIPTION	QTY	TOTAL	Delivered, Weeks/Months
(NTP).  52. The machine should have a US FDA Approval or CE Mark Approval. The bidder shall provide the manufacturer's approved US FDA Premarket Notification (PMN) or 510(k) certificate or CE Mark Approval certificate.  53. Certification that the supplier has the capability for corrective and preventive maintenance of the unit. The certificate must be issued by the manufacturer for the bidder.  54. Certification from the manufacturer authenticated by the Philippine consulate from the country of origin or Philippine branch of the manufacturer that the warranty should not be affected with a change of distributor.  55. Certification that the brand and model must be in the local market for a minimum of Five (5) years with at least five (5) Full digital Xray of the same brand in tertiary government facilities (at least 1 must be DOH hospitals).  56. Certification that the bidder is existing in the local market for a minimum of ten	ITEM NO.	Standard Requirements 47. The supplier shall secure a certificate from the manufacturer stating that the equipment is brand new, unused and not a discontinued model or was listed in the market recall. The certificate reflect the brand, model, serial number of the machine, and the date and place it was manufactured. 48. System must not be a retrofit solution Detector and x-ray machine are of the same brand. 49. The bidder must provide the current and valid certificate of Manufacturer's compliance with ISO 13485. 50. Supplier/ Bidder must be ISO certified 51. Delivery of equipment and all accessories within 90 calendar days upon receipt of Notice to Proceed (NTP). 52. The machine should have a US FDA Approval or CE Mark Approval. The bidder shall provide the manufacturer's approved US FDA Premarket Notification (PMN) or 510(k) certificate or CE Mark Approval certificate. 53. Certification that the supplier has the capability for corrective and preventive maintenance of the unit. The certificate must be issued by the manufacturer for the bidder. 54. Certification from the manufacturer authenticated by the Philippine consulate from the country of origin or Philippine branch of the manufacturer that the warranty should not be affected with a change of distributor. 55. Certification that the brand and model must be in the local market for a minimum of Five (5) years with at least five (5) Full digital Xray of the same brand in tertiary government facilities (at least 1 must be DOH hospitals). 56. Certification that the bidder is existing in		TOTAL	Weeks/Months

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	<ul> <li>57. Certification from the supplier will be responsible for the notification, transportation, delivery, installation, acceptance testing and commissioning and expenses for such will be on the account of the supplier.</li> <li>58. Must pass the performance/acceptance testing conducted by the FDA prior to acceptance by the end user</li> <li>59. Certification that the supplier/bidder shall provide applications training for users and maintenance personnel of the hospital.</li> <li>60. On-house training with certificate for the End-User Familiarization of the operating procedures of the equipment</li> </ul>			
	61. Certification to provide user's Manuals:  Two (2) sets of Service Manual in English Language upon delivery of the equipment  Two (2) sets of User Manual in English Language upon delivery of the equipment			
	<ul> <li>62. Certification that the suppliers/bidder shall provide free installation (includes electrical) of the equipment and its inclusions if needed.</li> <li>63. Current and Valid Certificate of Authorized or Exclusive Distributorship</li> <li>64. The manufacture must have an accredited service provider in the Philippines with parts available</li> <li>65.24/7 Technical Support.</li> <li>66. A certification of good performance from at least three (3) tertiary hospital stating the name of facility, address, type of equipment and its model, and the date of delivery must be provided.</li> </ul>			

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CONFORM (Signature	MED BY: e over printed name)		

DATE: