Section VII. Technical Specifications

INSTRUCTION: Bidders must state here either "Comply" or "Not Comply" against each of the individual parameters of each Specification stating the corresponding performance parameter of the equipment offered. Statements of "Comply" or "Not Comply" must be supported by evidence in a Bidders Bid and cross-referenced to that evidence. Evidence shall be in the form of manufacturer's un-amended sales literature, unconditional statements of specification and compliance issued by the manufacturer, samples, independent test data etc., as appropriate. A statement that is not supported by evidence or is subsequently found to be contradicted by the evidence presented will render the Bid under evaluation liable for rejection. A statement either in the Bidders statement of compliance or the supporting evidence that is found to be false either during Bid evaluation, post-qualification or the execution of the Contract may be regarded as fraudulent and render the Bidder or supplier liable for prosecution subject to the provisions of ITB Clause 3.1(a) (ii) and/or GCC Clause 2.1(a)(ii).

Public Bidding: One (1) Lot of Supply, Delivery, Installation, Testing, and Commissioning of Branded and Brand-new Magnetic Resonance Imaging (MRI) System inclusive of All Integral and Critical Components

Bidder's Proposed Brand Name:		Year & Model:	
Specifica- tion	Particular Requirements	Statement of Compliance	Supporting Documents for the Statement of Compliance (if applicable)
A. GENERAL REQUIRE- MENTS	1. The machine should be branded, brand new and a recent model (at least 2021)		
	2. The machine should be able to image the whole body.		
	3. The machine should be compatible and connected to our existing RISPACS with non-expiring Modality Work List License, at no additional cost to the Procuring Entity		
B. MAGNET SYSTEM	Superconducting Magnet: at least Tesla active shielded		
	2. Magnet length: 175 cm or below		
	3. Magnet bore: at least 60 cm		
	4. Shimming: active and passive		
	5. Magnet cooling (Cryogens): Helium (Zero boil off)		
	6. Magnetic field homogeneity: The magnetic field homogeneity should be equal to or less than:		
	• 45 cm DSV: 1.0 ppm		
	• 40 cm DSV: 0.45 ppm		
	• 30 cm DSV: 0.15 ppm		

	• 20 cm DSV: 0.06 ppm	
	• 10 cm DSV: 0.02 ppm	
C. GRADIENT SYSTEM	Maximum gradient amplitude (mT/m): At least 33 mT/m	
	2. Maximum slew rate (T/m/s): At least 100 T/m/s	
	3. Maximum Field of View (in xyz): 50x50x50 cm	
	4. Noise reduction:	
	With noise reduction	
	With vibration reduction	
D. RF SYS- TEM	• Water cooled	
	The RF computer (transmitter and receiver should be computer con- trolled and fully digital type	
	1. RF amplifier type: Solid state	
	2. RF Transmitter power: 10 kW or better	
	3. Number of Independent Digital RF receiving channels: At least 16	
E. RF COILS	Main body coil integrated to the magnet: Must be quadrature	
	Head-Neck Neurovascular Array: With at least 16 channels	
	3. Spine or Posterior Array embedded in the table: With at least 12 channels	
	4. Anterior Array: With at least 9 channels	
	5. Flex coil for shoulder, knee, and hip: With at least 8 channels	
	6. Flex coil for wrist and hand: With at least 8 Channels	
	7. Coil of foot and ankle: With at least 8 channels	
	8. Breast Array: With at least 8 channels (optional)	
	Should be fully motorized, computer controlled	
F. PATIENT TABLE	2. Movements in vertical and horizontal directions	
	3. Should be able to withstand patient load of 200 kg	

	4. With emergency switch for quick release of tabletop	
	With automatic sensing coil ports	
	6. Should have patient alarm	
G. MAIN COMPUTER	Main Operator Console Computer System	
CONSOLE AND WORK STATION	Manufacturer's specification	
(IMAGE PRO- CESSING	Multitasking capabilities (scan, film, and post processing) at the same time	
AND APPLI- CATION SOFTWARES	Simultaneous scan and recon	
	With two-way intercom system and Emergency stop button	
	• Image reconstruction speed of at least 60,000 FFT/second at 256x256 matrix	
	With CD/DVD writer	
	With noise and artifact reduction dur- ing reconstruction	
	With deep learning-based reconstruc- tion in all anatomies	
	2. Application Software	
	i. Imaging modes, Pulse Sequences, and capabilities	
	The following must be available with the MRI system:	
	• Spin echo (SE)	
	• Fast Spin Echo (FSE)	
	Gadient Echo (GE)	
	Spoiled Gradient Echo	
	Fast Steady State sequence	
	• FLAIR	
	• Fat and water saturation/separation	
	• 2D and 3D time of flight	
	• 2D and 3D phase contrast	

Non contrast MRCP	
Dual Echo Fast Gradient Echo	
Echo Planar Imaging (EPI)	
Short T1 Inversion Recovery (STIR)	
Double Inversion Recovery (DIR)	
Triple Inversion Recovery (TIR)	
• SWI	
Motion Correction	
• Spectroscopy	
Perfusion Study	
Tractography	
Diffusion Tensor Imaging	
Metal Artifact Reduction and Correction	
Compressed Sensing	
Whole body DWI	
• Enhanced Diffusion weighted imaging	
Multi shot Diffusion	
Respiratory gating/ trigger	
Deep Learning Reconstruction	
Volumetric imaging on T1, T2, FLAIR, PD	
Viability imaging	
Dynamic body imaging	
Dynamic brain function which in- cludes blood flow, blood volume, mean transit time, time to peak sig- nal intensity	

	Whole body metastasis screening	
	Non contrast vascular imaging	
	Fat suppression	
	Cerebral blood flow measurement	
	Non contrast MRA	
	3. MRI Workstation	
	 With three (3) sets of radiologist workstations with dual monitors and DICOM viewers with non-expiring li- censes, Windows operating software, Microsoft Office, and tables and chairs 	
	• Should be able to perform Neuro image processing like Perfusion, Diffusion, Perfusion Diffusion Mismatch, Spectroscopy	
	Should be able to perform body image processing	
H. OTHER	1. Accessories	
REQUIRE- MENTS	• One (1) MRI compatible CCTV system for patient observation	
	One (1) unit Music system for the patient	
	• One (1) unit Two-way intercom system	
	One (1) MRI compatible patient trolley	
	One (1) MRI compatible wheelchair	
	Two (2) MRI compatible IV stand	
	One (1) MRI compatible coil cabinet	
	Two (2) Hand held metal detectors	
	One (1) Laser printer or photopaper or dry film printer	
	One (1) heavy duty printer and scanner with refillable ink tank	
	• Two (2) MRI compatible Fire Extinguishers	
	One (1) MRI compatible stretcher/gurney	

One (1) MRI Compatible transfer board		
Two (2) MRI compatible medical oxygen regulator/guage		
• Two (2) MRI compatible oxygen tanks (at least 40 liters		
One (1) Oxygen monitor in the Magnet room		
One thousand pieces (1000) ear plugs		
One thousand pieces (1000) Head set covers		
One (1) MRI compatible suction machine (optional)		
• One (1) set MRI phantom		
• Two (2) units air purifiers with at least 500m3/h CADR and H13 filters with ten (10) extra H13 filters		
With appropriate MR cooling/ chiller system		
Security straps, warning signs, and positioning aids		
Fifteen (15) pcs Gadobutrol 5 ml contrasts		
One (1) unit compatible pressure injector		
• Thirty (30) units each of syringes, tubings, contrast bottles compatible with the pressure injector		
• Three (3) units of UV light, at least 58W, non-ozone (optional)		
One (1) unit MRI compatible pulse oximeter		
• One (1) unit Network attached storage, at least 100 TB storage and 5 bays		
2. Airconditioning		
• Two (2) units appropriate air conditioning system for the magnet room		
Two (2) units appropriate air conditioning system for the equipment room		
• One (1) unit appropriate air conditioning system for the control room		
One (1) unit appropriate air conditioning system for the recovery/preparation room		
One (1) unit appropriate air conditioning system for the waiting area		

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Two (2) units appropriate air conditioning system—for the reading and doctor's room	
One (1) set of appropriate dehumidifier	
All conditioning units must be inverter type	
3. MRI Room preparation	
• RF shielding for the MRI room should be provided.	
• Magnetic shielding should also be provided if needed.	
4. Civil works and Electrical works	
• The needed civil and electrical works are included (Plans are done by Engi- neering Section)	
Step up or step down transformer if needed	
• The supplier must provide connection to the hospital's main power line	
One (1) unit UPS for the MRI scanner and chiller with at least 10 minutes backup time	
• One (1) Transient voltage surge sup- pressor (TVSS) compatible with the MRI Scanner	
5. Warranty, Maintenance, Service	
• Three (3) years comprehensive war- ranty on parts and services of the MRI machine to start after end users' ac- ceptance	
At least one (1) calibration per year	
At least four (4) preventive mainte- nance per year	
6. Trainings	
At least two (2) weeks or 80 hours off site training for 2 Radiologic Technologists to a health facility with the same MRI model at no cost to the Procuring Entity	
• At least two (2) weeks or 80 hours on site training for 2 Radiologic Tech- nologists, Radiologists, and Physicist at no cost to the Procuring Entity	
The Supplier shall provide training certificates upon completion of all training requirements	
7. Others	
• Product data sheet of the MRI in English should be submitted as attachment to support the Statement of Conformity to Section VII Technical Specifica-	
tions.	

	Must pass the performance/ acceptance testing conducted by FDA or its accredited testing body	
	Must pass the requirements for FDA registration/ licensing	
	• Undertaking to provide magnetic shielding, if necessary for the projects upon conduct of site inspection.	
	Certification that the manufacturer has been in the business of manufac- turing MRIs for at least 20 years	
	Certification that the brand being offered has at least 15 MRI installations in the Philippines in the last five (5) years and can be visited at End user's disposal	
	Certification that the brand that being offered has been in the local market for at least 15 years	
	Certification by the manufacturer that supplies, parts, and accessories shall be available for at least 15 years after expiration of warranty	
	Certification that service engineers trained on service and repair are available in the Philippines	
	Certification that the manufacturer has a local office in the Philippines for the last 15 years with SEC registration	
	MRI machine should be US FDA approved	
	Certification of Manufacturer's ISO compliance to 13485:2016	
	Certification that the MRI machine proposed is a current model launched within the past 3 years (at least 2021)	
	Service and operations manuals, soft and hard copies	
I. TURN KEY	Room renovation and other civil works including floor, wall, ceiling finish, appropriate doors, and furni- ture in the MRI area	
	Electrical works within the MRI area	
	Plans and terms of references for the MRI area including civil works, elec- trical works, furniture are care of En- gineering Section	
CONFORME:		
(Company Name)		
(Name and Signature of Authorized Representative)		