

Republic of the Philippines
Department of Budget and Management
PROCUREMENT SERVICE PHILIPPINE GOVERNMENT ELECTRONIC PROCUREMENT SYSTEM
BIDS AND AWARDS COMMITTEE



Supplemental/ Bid Bulletin No. 6 22 July 2019

PUBLIC BIDDING No. 19 - 142 - 4

Supply, Delivery, Installation, Testing, and Commissioning of Nuclear Medical Equipment and Room Shielding for the Philippine Orthopedic Center

Issued pursuant to Sec. 22.5 of the IRR of R.A. 9184 to clarify and/or amend certain provisions in the Bidding Documents issued for this project, considering the issues raised and clarifications made by prospective bidders during the Pre-Bid Conference held on **July 4, 2019**, likewise, respond to bidders' written queries received within the prescriptive period for filing.

I. Amendments

REFERENCE	BASES FOR AMENDMENT / INCLUSION
SECTION VI. Schedule of Requirements Page 69 Within One Hundred Five (105) Eighty (180) Calendar Days from receipt of Notice to Proceed including the Delivery, Installation, Testing, and, Commissioning.	Refer to the revised Schedule of Requirements attached as Attachment "A".
Section VII. Technical Specifications xxx 6. ACQUISITION SYSTEM REQUIREMENTS xxx e. Energy window width of 160kEV-up to 600keV at least 35 keV up to 588 KeV or higher xxx	Refer to the revised Technical Specifications attached as Attachment "B".
Section VII. Technical Specifications xxx 14. QUALITY CONTROL xxx i. Thyroid Phantom xxx	Refer to the revised Technical Specifications attached as Attachment "B".



For the purpose of this Bulletin and for better understanding of its contents, the following rules shall apply: (a)

Bouble Strike out — denotes deletion; (b) Underline — denotes inclusion of new item/requirement, and "xxx"

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Section VII. Technical Specifications

XXX

9. WHOLE BODY ACQUISITION

Whole body scan length: 200 cm maximum length <u>or</u> <u>higher</u>

XXX

Refer to the revised Technical Specifications attached as Attachment "B".

Section VII. Technical Specifications

XXX

11. SPECT ACQUISITION

a. SPECT with step and shoot and acquire during step acquisition — Variable zoom factors $\frac{\text{up-to}}{\text{of}}$ 3.0 or greater

XXX

Refer to the revised Technical Specifications attached as Attachment "B".

II. Clarifications

ITEM NO.	ISSUE/ REQUEST	CLARIFICATION/ RESOLUTION				
	ASSURANCE CONTROLS TECHNOLOGIES CO., INC.1					
	SECTION VI. Schedule of Requirements Page 69 Within One Hundred Five (105) Calendar Days from receipt of Notice to Proceed including the Delivery, Installation, Testing, and Commissioning.	Please refer to the discussion in I. Amendment.				
	Normally, the infrastructure would take 3-4 months to finish. Since the site is not yet ready to accommodate the delivery and installation by the time of receipt of the Notice to Proceed, it would be best to adjust the delivery period to allow the on time delivery of the equipment and installation.					

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¹ Received on 19 July 2019

Section VII. Technical Specifications The winning bidder will only provide Room Shielding. The air-conditioning, Shielding -not less than 6 feet height with at architectural, plumbing, and electrical least 1.5 mm lead sheet thickness or works will be the responsibility of the Endequivalence in concrete, applicable to lead doors User. and walls as necessary. XXX Please identify the Scope of Works for the Project. We would like to clarify if the airconditioning, civil, architectural, plumbing and electrical works are included in the scope of works as these are vital in the costing for the whole project. Section VII. Technical Specifications Please refer to the discussion in I. Amendment. 6. ACQUISITION SYSTEM REQUIREMENTS e. Energy window width up to 160keV up to 600 keV XXX The range of 160keV up to 600keV favors only 1 supplier. Our energy window width range is 35keV up to 588keV. Adjusting the range would open up the chance to other suppliers who wish to join the bidding. **Section VII. Technical Specifications** Please refer to the discussion in I. Amendment. 14. QUALITY CONTROL XXX i. Thyroid Phantom XXX The item is no longer available in the market.



Section VII. Technical Specifications Please refer to the discussion in I. Amendment. III. SPECIFICATION FOR UPTAKE MACHINE & HOT LABORATORY 1. Thyroid Uptake System with Well Counter b. Detector: 3/8" Shielded Nal (Tl) crystal XXX No available literature indicating dimensions. Appropriate and compatible Shielded Nal (TI) crystal with the Uptake Machine shall be provided. **Section VII. Technical Specifications** Please refer to the discussion in I. Amendment. c. Multichannel Analyzer: 1800 Channels What we have is 1024 Multichannel Analyzer Channels. 1800 Channels is only available to one brand. **Section VII. Technical Specifications** Please refer to the discussion in I. Amendment. 2. Neck Phantom for Thyroid Uptake d. I.D.: 4"h x 2" diameter (10 x 5 cm) XXX For clarification: What does "I.D." mean? We would like to request its removal. Usually, the description is used by only one brand. Section VII. Technical Specifications Please refer to the discussion in I. XXX Amendment. 3. Digital Dose Calibrator c. Automatic range selection up to 100 Curies of Tc 99 or 25 Curies of F-18 8 Curies of Tc 99 is enough since PNRI does not give license to possess higher than 8 Curies, high activity Tc99m are measured in dispensing radiopharmacies. Meanwhile, the standard range of F-18 is only 20 Curies. The range selection of 100 Curies Tc 99 and 25 Curies of F-18 is only available in one brand.

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Please refer to the discussion in I. Section VII. Technical Specifications Amendment. XXX 7.17 One (1) Movable Lead Barrier with Lead Plastic Window c. Four hospital grade: two locking and two nonlocking XXX For clarification if it's "four" or "for" hospital grade. Does locking and non-locking refer to the caster wheels? **Section VII. Technical Specifications** Please refer to the discussion in I. Amendment. 4. PATIENT BED f. Patient Table: Maximum patient load shall be less than or equal to 220 kgs For clarification: Since this is about patient load, will a higher maximum patient load be acceptable? Our maximum patient load capacity is 227kgs (500lb). Section VII. Technical Specifications Please refer to the discussion in I. Amendment. 9. WHOLE BODY ACQUISITION Whole body scan length; 200 cm maximum length XXX For clarification: Will it be acceptable if our maximum length would be greater than the requirement? Ours is 203cm and a 3cm difference would not limit you to scan longer patients. Please refer to the discussion in I. **Section VII. Technical Specifications** Amendment. XXX 11. SPECT ACQUISITION a. SPECT with step and shoot and acquire during step acquisition - Variable zoom up to 3.0 or greater XXX It would be confusing to retain the word "up to" since the required variable zoom is greater than or equal to 3.0.

For the purpose of this Bulletin and for better understanding of its contents, the following rules shall apply: (a) Double Strike out — denotes deletion; (b) Underline — denotes inclusion or new item/requirement; and "xxx" — denotes separation of phrase/s being amended from the rest of the main text.

a. Three (3) months' supply of free radiopharmaceuticals including radio-active iodine 131 for therapy	The Technical Specifications provides for the minimum requirement and the prospective bidder is not precluded from providing higher offers.
Assurance Technologies is suggesting to the above requirement stating that it will be beneficial for the calibration if the machine and test-patients.	
b. Free of charge licensing assistance (PNRI and DOH) Assurance Technologies is suggesting to add the above requirement stating that it will help the facility to boost their start-up operation and productivity.	Please refer to Bid Bulletin No. 4. Where a Certification that upon receipt of the Notice to Proceed the bidder will apply for licensing with the Philippine Nuclear Research Institute prior to the installation and commissioning of the Nuclear Medical Equipment will be required.

The herein amendments form an integral part of the bidding documents. Correspondingly, all other provisions in the bidding documents affected by these amendments are similarly amended or modified.

The clarifications made, explain in greater detail the purpose or intent of the requirement and do not necessarily amend that particular provision in the bidding documents.

(SGD.) ENGR. ESTRELLITA G. FULE Chairperson, BAC IV

Schedule of Requirements

Lot No.	Item/ Description	Quantity	Delivery Period
1	Supply, Delivery, Installation, Testing, and Commissioning of Nuclear Medical Equipment and Room Shielding for the Philippine Orthopedic Center	1 Lot	Within One Hundred Eighty (180) Calendar Days from receipt of Notice to Proceed including the Delivery, Installation, Testing, and Commissioning.

I hereby commit to comply and deliver all the above requirements in accordance with the above stated schedule.

Name of Company	Signature Over Printed Name of	Date
	Authorized Representative	

Technical Specifications

LOT NO. 1		Supply, Delivery, Installation, Testing, and Commissioning of Nuclear Medical Equipment and Room Shielding for the Philippine Orthopedic Center
QUANTITY	:	One (1) Lot
APPROVED BUDGET FOR THE CONTRACT	:	¥ 52,000,000.00

AGENCY SPECIFICATIONS	BIDDER'S STATEMENT OF COMPLIANCE	ACTUAL OFFER	REFERENCE
Conforms with the following minimum requirements:			
A. Dual Head Gamma Camera with		Brand and Model:	
Single-Photon Emission Computed			
Tomography-Computed			
Tomography (SPECT/CT Scan)			
Shielding –not less than 6 feet in			
height with at least 1.5 mm lead			
sheet thickness or equivalence in			
concrete, applicable to lead doors			
and walls as necessary.			
Shielding should be in place in the			
following rooms:			
SPECT/CT Room,			
Uptake Room,			
Bone Densitometer Room,			
Hot Laboratory Room,			
Decontamination Room,			
Radioisotope Storage Room,			
Post Admin Room,			
Dose Admin Room, and			
Radioactive Patient Toilet			
	·		
External partition walls will be			
provided by the End-User.			
I. GAMMA DETECTOR			
a. Detectors shall be shielded for			
high energy range of 250 keV or			
higher	-		
b. Number of Detectors: Two (2)			<u></u>

AGENCY SPECIFICATIONS	BIDDER'S STATEMENT OF COMPLIANCE	ACTUAL OFFER	REFERENCE
c. With true rectangular Field of			
view (FOV)(i.e., FOV corners not			
clipped for wider FOV and better	**		·. ·
appreciation of images) d. Field of View shall be equal or			
larger than 52 cm x 37 cm (20.5 in x			
14.5 in) Crystal Thickness			
e. Number of PMTs /Detector > 56			
II. NEMA SPECIFICATIONS (Minimum			
Requirements using the appropriate			
NEMA Standards)			
-A. Intrinsic Spatial Resolution (typical)	·		
a. FWHM for CFOV < 4.0 mm			
b. FWHM for UFOV < 4.0 mm			
c. FWTM for CFOV < 8.0 mm			
d. FWTM for UFOV < 8.0 mm			
B. Intrinsic Spatial Linearity			
a. Differential CFOV < 0.25 mm			
b. Differential UFOV < 0.25 mm			
c. Absolute CFOV < 0.5 mm			
d. Absolute UFOV < 0.8 mm			
C. Maximum count rate (per detector) > 300 000 cps			
D. System Sensitivity per detector			
(Tc-99m, LEHR collimator) >			
160cts/min/uCi 1. GANTRY	·		
a. The gantry should support variable angle configurability of the			
detectors including 90°, 180° SPECT,			
and other angles useful for SPECT.			
b. At least one of the detectors shall			
permit caudal and cephalic			
tiltgreater than or equal to 15	·		
degress or equivalent, allowing			
detector positioning close to imaging area and detector motion shall allow			
patient imaging in sitting and			
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	BIDDER'S		
AGENCY SPECIFICATIONS	STATEMENT OF COMPLIANCE	ACTUAL OFFER	REFERENCE
standing positions.			
c. The system shall support Step			
and Shoot and Continuous SPECT			
detector rotation modes.			
d. The system shall support Non-			
circular orbits and automatic			
contouring for SPECT Acquisitions			
with all detector configurations (90°			
and 180°)			
e. The gantry shall have an opening			
of at least 70 cm			
f. Necessary hand controls, for gantry and detector motion, shall be			
provided on both sides of the gantry.			
g. The gantry shall have safety			
features including emergency stop			
buttons on both sides of the gantry			
and patient contact sensors on each			
collimator			
h. The gantry shall be linked to the			
patient table and have the necessary			
sensors to recognize the patient			
table position at all times to prevent			
accidental collisions.			
i. The system shall be able to			
perform non-uniform attenuation			
correction using CT Attenuation			
maps acquired in the same system,			
for general SPECT imaging. 2. GANTRY AND ACQUISITION			
STATUS			
Patient positioning monitor (PPM) at			
the gantry display monitor shows	,		
status of the acquisition.			ļ
3. SPECT/CT FEATURES &			
CAPABILITIES			
a. CT can acquire at least two (2)			
slices or better, interleaved			
reconstruction per rotation		-	
b. Minimum CT Slice Thickness: <			
1mm			
c. The CT scan required for			
attenuation correction and			<u> </u>

	BIDDER'S		
AGENCY SPECIFICATIONS	STATEMENT OF COMPLIANCE	ACTUAL OFFER	REFERENCE
anatomical mapping shall not add	,		
more than 30 seconds to the total			
SPECT/CT acquisition time.			
d. The system shall be capable of			
automatically matching the CT slice			
thickness to the SPECT slice			
thickness for accurate image fusion			
and attenuation correction			
e. CT Field of View Diameter: 70cm			
f. The system shall offer a	•		
technology that reduces the			
unnecessary CT dose			•
g. Gantry Port Diameter: 70cm			
h. Continuous spiral CT range should			
be greater than or equal to 150 cm			
i. Tube Anode heat storage			
capacity: 2.0 MHU and above			
j. Tube Current up to 200mA or			
higher			
k. Selection of Tube voltage up to 130 kV			
I. Reconstructed slice width of 1	- 1- 1- 1- 1- 1- 1- 1- 1- 1- 1- 1- 1- 1-		
mm			
m. Scan times for full 360 degree	-		
scan of 0.98s or faster			
n. High contrast resolution at 0%			
MTF (+/-10%)should be 15 lp/cm or			
higher			
4. PATIENT BED			
a. With motorized vertical and			
horizontal motion activated from the			
hand controls and preset positions.			·
b. Minimum Patient bed height:			
55cm			
c. Patient bed shall have ability to			
position any part of body under the			
detectors without moving the			
patient. All pallet motions shall be activated from the hand controller.			
d. The patient bed shall have < 10 %		<u> </u>	
attenuation for 140 keV photons.			
e. Whole body scan Length shall be			1
c. Whole body scall Lelight shall be		<u> </u>	<u> </u>

AGENCY SPECIFICATIONS	BIDDER'S STATEMENT OF COMPLIANCE	ACTUAL OFFER	REFERENCE
up to 200 cm			
f. Patient Table: Maximum patient			
load shall be 220 kgs or higher			
g. ECG Cable port integrated into			
Bed or Gantry			
5. COMPUTER SYSTEM MINIMUM REQUIREMENTS			
a. Acquisition Workplace section:			
Customizable Display			
b. Acquisition Workplace section:			
Customizable Workflows			
c. Two (2) workstations (WS): 1WS			
for acquisition and 1WS for post-			
processing and reading			
d. All organ processing software			
(renal, lungs, bone, GIT, liver and			
neuro protocols) e. Appropriate and Authentic			
Licenses for operating system			
software.			
f. Conversion data files to DICOM			
format integrated to existing	;		
hospital information system and			
modality worklist.			
6. ACQUISITION SYSTEM REQUIREMENTS			
a. User shall have the ability to			
modify acquisition parameters easily			
and quickly.			
b. Simultaneous acquisition and			
processing capability on same	i		
computer			
c. Independent energy window			
selection			
d. Number of energy windows			
supported should be at least 6 windows per detector			
e. Energy window width of at least		-	
35 keV up to 588 keV or higher			
f. The system shall support			
symmetric and asymmetric energy			
windows			
g. The system shall offer manual			
8. The system shall offer mariant		<u> </u>	•

AGENCY SPECIFICATIONS	BIDDER'S STATEMENT OF COMPLIANCE	ACTUAL OFFER	REFERENCE
and automatic annotation (patient,			
study)			
h. Start and stop acquisition control from:			·
i. Camera hand control			
il. Computer			
i. Allow the user to combine			
acquisition and processing of protocols in one protocol			
j. Capable of combining multiple			
SPECT acquisitions (e.g. Cardiac			
Stress & Rest acquisitions) in one protocol.			
k. ECG compatible to the system			
shall be provided and connected.			
l. Acquire cardiac data in-half the			
time (half-time imaging)			
7. STATIC ACQUISITION			
Matrix size			
a. 64 x 64			
b. 128 x 128			
c. 512 x 512			
d. 1024 x1024			
8. DYNAMIC IMAGE ACQUISITION			
Matrix size			
a. 64 x 64			
b. 128 x 128			
c. 256 x 256			
9. WHOLE BODY ACQUISITION			
Whole body scan length: 200 cm or higher			
10.GATED IMAGE ACQUISITION			
Matrix Sizes			
a. 64 x 64			
b. 128 x 128			
c. Buffered beat			
d. Bad beat rejection			

AGENCY SPECIFICATIONS	BIDDER'S STATEMENT OF COMPLIANCE	ACTUAL OFFER	REFERENCE
11.SPECT ACQUISITION			
a. SPECT with step and shoot and acquire during step acquisition - Variable zoom factors of 3.0 or greater			
b. Variable Start angle		·	
c. Dual isotope SPECT capability			
12.GATED SPECT ACQUISITION			
a. Matrix Sizes			
i. 64 x 64			
ii. 128 x 128			·
b. Buffered beat			
c. Accepted and rejected beats shall be saved separately in the patient file to ensure high statistical accuracy with the summed image d. Forward/Backward framing by a			
user-defined percentage			
e. End study by time per view or number of accepted beats per view			
13.COLLIMATORS			
 a. Collimators change should include some level of automation. b. Collimator changing shall be possible without moving the patient 			
table or partial movement of table			
c. Low Energy High Resolution			
d. High Energy Collimator			
e. Pinhole Collimator			
f. General Purpose			
14.QUALITY CONTROL			
a. Integrated / Supplied Source Holder for QC			
b. Simultaneous QC for Both Detectors			
c. Energy Independent QC	-		
d. Four-quadrant bar phantom			

	BIDDER'S		
AGENCY SPECIFICATIONS	STATEMENT OF COMPLIANCE	ACTUAL OFFER	REFERENCE
e. Flood Phantom for Technetium			
f. Flood Phantom for Iodine			
g. Linearity Phantom			·
h. ECT phantom			
15. INSTALLATION REQUIREMENTS			
a. One (1) Uninterruptible Power Supply (UPS) for the Workstations			
- Should be compatible with the workstation			
b. One (1) Uninterruptible Power Supply (UPS) for the Gamma Camera			
- Should be compatible with the wattage of the Gamma Camera SPECT/CT			
c. Transformer and TVSS for the CT			
- Should be compatible with the wattage of the Gamma Camera SPECT/CT and UPS			
d. Lead Glass: 2.1mm Pb, 100 x 120cm			
III. SPECIFICATION FOR UPTAKE MACHINE & HOT LABORATORY			
THYROID UPTAKE SYSTEM WITH WELL COUNTER - RADIOACTIVE IODINE UPTAKE AND THERAPY a. 15" VGA LCD Colored Screen			
Display or higher			
b. Detector: appropriate shielded Nal (TI) crystal			
c. Multichannel Analyzer: 1024 Channels or higher; Automatic self- calibration of 2000 counts per			
seconds;Intrinsic EnergyResolution of (IER) 9.9%; Intrinsic Count Rate Performance of 280 CPS; Multiple			
Window Spatial Registration of 1.0mm; Energy Range of 40KeV-			
300KeV; Correction of para-injection and empty syringe. ROI - Automatic or Manual;			
Background subtraction and mean			

BIDDER'S			
AGENCY SPECIFICATIONS	STATEMENT OF COMPLIANCE	ACTUAL OFFER	REFERENCE
energy calculation via curve fittings;			
with ; Linear Display -			
Automatic/Manual			
d. Nuclide Data: Over 90 Nuclides in			
Memory (major gamma-ray		•	
energies, keV and half-life)			
e. Advanced System Setup: Test			
sources, Efficiencies, User Nuclides,			
Bioassay Data, Thyroid Uptake			
protocols, Thyroid Uptake Normal			
Values			
f. Diagnostics and Tests: Full system			
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Į.			
d. 4"h x 2"diameter (10 x 5 cm)			
e. Should include bottle carrier,			
capsule holder and 12 polyethylene			
bottles			
3. DIGITAL DOSE CALIBRATOR			
· · · · · · · · · · · · · · · · · · ·	drasa		
· · ·			
management system.			
c. Automatic range selection up to			
8 Curies of Tc 99 or 20 Curies of F-18			
self-diagnostics including all program and data memories; Comprehensive test programs include automatic Chi-Square, MDA and FWHM g. Printer: Color Inkjet Resolution: 1200 x 1200 dpi Power Specifications: 230 volts 2. NECK PHANTOM FOR THYROID UPTAKE a. Made of clear lucite Poly (methyl methacrylate) b. With two (2) part insert for bottle counting and vial capsule counting c. Phantom Dimensions: 5"h x 5" diameter (127 x 127 cm) d. 4"h x 2"diameter (10 x 5 cm) e. Should include bottle carrier, capsule holder and 12 polyethylene bottles 3. DIGITAL DOSE CALIBRATOR Combination of the Dose Calibrator and Well Counter 3.1.SPECIFICATIONS a. Display Screen: touch screen display b. Communications: Ethernet, serial port for Nuclear medicine management system. c. Automatic range selection up to			

AGENCY SPECIFICATIONS	BIDDER'S STATEMENT OF COMPLIANCE	ACTUAL OFFER	REFERENCE
d. Display in Curies or Becquerels			
e. Library of over 80 nuclides with calibration number and half-life and room for 10 additional nuclides			
f. Over 80 Nuclides with half-lives in memory			
g. 64 Channel MCA			
h. Built-in dose calibration, quality control and self-diagnostics to ensure longer life accuracy			
i. Automated QC including constancy and linearity programs			
j. USB/PC Communications			
k. Software upgrade via Ethernet interface			
I. High sensitivity, drilled NaI well crystal			
m. Energy Spectrum: 0-800 Kev			
n. Wipe test result and QA data can be stored in memory and printed at any time			
o. Measurement Range: Auto- ranging, up to 250GBq			
p. Energy Range: 20KeV			
q. Format: Direct reading in Bq or Ci — User selectable or fixed			
r. Response Time: one or two seconds for doses greater than 200uCi, three seconds for doses greater than 20 uCi; 50-100 seconds below 20 uCi of Tc-99m with default threshold			
s. Electrometer: Accuracy: +-1% or 0.2 uCi whgichever is greater			
t. Repeatability: Within +-0.3% above 1 mCi short term in 24 hours and 1% long term in 1 year			
3.2.TESTS: DIAGNOSTICS			
a. Full test of program, system memories; Daily – Auto Zero, Background Adjust, Data Check,			

AGENCY SPECIFICATIONS	BIDDER'S STATEMENT OF COMPLIANCE	ACTUAL OFFER	REFERENCE
Accuracy and Constancy, Enhanced – Linearity, Geometry, Strip QC	10 and 10		
3.3.NUCLEAR DATA			
a. Nuclide Keys – 10 preset keys keys; System Memory – Over 80 nuclides (cal number and half-life)			
4. RADIOIODINE FUME HOOD			
a. Dosage Cabinet with stainless steel frame on both sides and glass door in front.			
b. 0.60mW x 0.72mD x 1.2 mH			
c. with Charcoal Filter			
5. LAMINAR FLOW HOOD			
 a. Working area:1200 x 600 x 600mm (w x d x h) b. Internal and external cabinet: stainless steel sheet c. Absolute filter for air outlet — 			
HEPA-H14			
6. UNIVERSAL POWER SUPPLY			
a. Should be compatible with the Laminar Flow Hood			
b. At least 1KVA			
7. ACCESSORIES			
7.1 One(1) Survey Meter calibrated for appropriate radio nuclei a. Alpha, beta, gamma and x-ray			
detection b. Multiplier Ranges: x0.1; x1; x 10; 100 for external detector; 1000 for			
internal detector c. Meter Face: 0-2 mR/hr, 0-2			
mR/hr, 0-6.6k cpm d. Reset switch: Push button to zero meter after over range exposure			
e. Detectors: Internal — Energy compensated GM, for high range gamma detection only; 2000mR/hr			
f. Detector: Pancake type halogen quenched GM			

AGENCY SPECIFICATIONS	BIDDER'S STATEMENT OF COMPLIANCE	ACTUAL OFFER	REFERENCE
g. Meter Face Dimension: 6.1 x 3.6			
h. Probe Holder: Unique tongue and		5 Page 24 To 10 A 10	
grove probe holder for one or two			
handed surface mounting.			
I. Sensitivity: 2100 cpm/mR/hr for Cs-137 dose			
j. Batteries: Two each, size "D", typical life 600 hours	W-14-		
7.2One (1) Contamination Meter			
calibrated for appropriate radio			
a. Three range surface rate meter			
with 2" built-in diameter pancake			
gauge memory detector			
b. Read-out is in counts per minute (and mR/hr).			~
c. Ranges: Linear – 0-500, 0-5,000, 0-50,000cpm			
d. Switch Position: Off, Battery Test, x100, x10, x1			
e. Audio: Internally mounted speaker			
f. Detector: Halogen-quenched "Pancake GM Tube"		,	
g. Diameter: 2" (5cm)			
h. Window Thickness: 1.5mg/cm2			
i. Background: Typical 50cmp. Thin profile of tube (13mm) gives low background			
j. Efficiency: 100% for all betas and alphas that have energy to penetrate the thin window			
k. Voltage: 900V Nominal			
I. Gamma Sensitivity: Nominal is 3000 cpm/mr/h (based on Cs-137)			
m. Feet: Replaceable neoprene feet			
n. Calibration: Single master calibration potentiometer and individual calibration pots for each scale.			
o. Power: 9 volt nominal "transistor			

AGENCY SPECIFICATIONS	BIDDER'S STATEMENT OF COMPLIANCE	ACTUAL OFFER	REFERENCE
battery" mercury or equivalent			
p. Current Drain: 3 mA typical			
q. Handle: Swivel Type polished anodized aluminum			
r. Battery Life: 100 hours in normal operation.			
7.3Co-57, Cs-137 & Ba-133 Reference Standard for Dose Calibrator			
7.4Personal Radiation Protection			
a. Three(3) Lead aprons with light weight flexible Lead vinyl with 0.5 mm lead attenuation			
b. Three (3) thyroid shields with light weight flexible Lead vinyl with 0.5 mm lead attenuation			
c. Two (2) pairs of lead glove			
c.1) Protective Gloves for x-ray			
c.2) 0.5mm lead equivalence			
d. Two (2) pairs of lead goggles	•		
d.1) 2" x 4.25" single sheet of			
fluoroscopic quality lead glass d.2) Glass provides 2.00 mm lead equivalency			
e. Two (2) Direct Read Dosimeter			
e.1) Range: 0-200mR			
f. One (1) Dosimeter Charger			
f.1 Capable of charging any Direct- Reading Dosimeter			·
f.2 Conforms to ANSI N42.6-1980			
f.3 Controls: One-Turn Potentiometer			
f.4 Reading: Spring-Loaded Push Rod	,	,	
f.5 Operating Temperature: 0-120F (-18-49C);			
f.6 Lamp: LED			
7.5 One (1) Benchtop Clear Lead "L" Shield			
a. Dimension: maximum of 281.94 mm w x 281.94 mm			

	BIDDER'S		
AGENCY SPECIFICATIONS	STATEMENT OF COMPLIANCE	ACTUAL OFFER	REFERENCE
b. Lead Shielding: 5" thick (1.2 cm)			
c. Viewing Panel: 2mm lead lead equivalent			
d. Clear plexi glass for Beta Shielding			
7.6 One(1) Benchtop Clear Lead "L" Shield			
a. Dimensions of 281 mm high x 281 mm viewing area			
b. Lead Equivalent: 2mm			
7.7 Two (2) 3cc Tungsten Syringe Shields			
a. Barrel shield with 2 mm thick tungsten			
b. With 5.0 density lead window			
c. With reflective internal surface for	•		
easy reading of the syringe markings			
7.8 Two (2) 5cc Tungsten Syringe Shields			
a. Barrel shield with 2 mm thick tungsten			
b. With 5.0 density lead window			
c. With reflective internal surface for easy reading of the syringe markings			
7.9 One (1) Small Shielded Waste Bin			
a. 10 to 20 mm lead shielding			
b. 7 to 12 liter capacity with pedal or handle cover			
7.10 One (1) Large Shielded Waste Bin			
a. 10 to 20 mm lead shielding			'
b. With 14 to 20 liter capacity with pedal or handle cover			
7.11 One (1) Niptong			
a. Specific for Nuclear Medicine Use			
7.12 One (1) Forceps			
a. Specific for Nuclear Medicine Use			
7.13 One (1) Set Rectangular interlocking Lead Bricks			

	BIDDER'S		
AGENCY SPECIFICATIONS	STATEMENT OF COMPLIANCE	ACTUAL OFFER	REFERENCE
a. Depend on the size of dose			
calibrator shield			
b. With V-shaped edges and			
common straight-edge bricks			
interlocking to cover the area of the	·		
dose calibrator shield			-
7.14 One (1) Decontamination Kit which contains the following:			
a. One (1) 30 gallon fiber drum		P-1 A-1-	
b. Two (2) pairs of coverall,		<u> </u>	
disposable			
c. Two (2) pairs Shoe cover,			
disposable			
d. Two (2) disposable nostril type			
Respirators			
e. Four (4) pieces eight inches by			
eleven inches (8" x 11") size Filters			
f. Two (2) Pairs Gloves, reusable			
g. One (1) gallon Radiation			
Decontamination Wash			
h. One (1) canister Radiation Decontamination Wipes			
i. One (1) bottle of One (1) Liter			
Radiation Decontamination Spray			
Mist			
j. Ten (10) Poly bags at least 6" x 9"			
x 2 mil			
k. One (1) piece metal 12" Niptong			-
I. One (1) piece hand Sponge			
m.One (1) piece standard Mop			
n. One (1) piece hand Scrub Brush			
o. One (1) piece 5 liter pail			
p. One (1) piece 5 meter at least 9.0			
mm thickness rope			
q. One (1) set of at least 5 pieces of the following:	·		
q.1) radiation danger warning			
q.2) radiation contamination sign			
q.3) Emergency sign with radiation hazard.			4

AGENCY SPECIFICATIONS	BIDDER'S STATEMENT OF	ACTUAL OFFER	REFERENCE
	COMPLIANCE		
7.15 Radiation Warning Signs and			
Labels			
a. For small room: 210mm x 85			
mm (at least 10 pieces)			
b. For large room: 280 mm x 122			
mm (at least 5 pieces)		-	
7.16 One (1) Leaded Sharps Container			
a. Compatible for Nuclear Medicine			
Waste			
b. With lockable hinge access			
7.17 One (1) Movable Lead Barrier			
with Lead Plastic Window			
a. Opaque Panel with 0.8mm Lead			1 1 1114 114
Casters			
b. Shielding Window: 0.5 mm lead			
equivalency			
c. Four hospital grade caster			
wheels: two locking and two non-			
locking			
7.18 One (1) Laboratory Cart, Stainless Steel			
a. Stainless steel			
b. With four (4) wheels			
7.19 One (1) Dehumidifier			·
a. Water Container Capacity:			
Minimum of five (5) liters			
7.20 One (1) Temperature &			
Humidity Monitor (For Gamma			
Camera room)			
a. Indoor monitor with temperature			
range of 0 to 50°C			
b. Humidity range of 16% to 98%			
7.21 One (1) Moly Assay Canister			
a. 7.6 mm Lead Shielding			
b. Standard size			
7.22 One (1) Elution Vial Shield			
a. Lead glass thickness: 14 mm			
b. Shielding thickness: 6mm lead			
with 360 view point	<u> </u>		•

	BIDDER'S		
AGENCY SPECIFICATIONS	STATEMENT OF COMPLIANCE	ACTUAL OFFER	REFERENCE
7.23 Radioaerosol Administration			
System for V-Q Scan enclosure is			
lead-shielded from top to bottom			
with oxygen dedicated external port			
and nebulizer attachments			
7.24 One (1) Urea Breath Test			
(14Carbon) starter kit set			
B. DEXA CENTRAL DUAL ENERGY X-			
RAY ABSORPTIOMETRY (DEXA) BONE			
DENSITOMETER			
a. Scanning method : Linear x-ray			
fan-beam with motorized table and			
motorized C-arm.			
b. Detector system: High density			
multi-detector array assembly.			
c. X-ray system: Dual-energy			· ·
100kVp/140kVp			
d. With automated internal			
calibration system and capable of			
storing and analyzing data			
e. Single energy scan switch capability			
f. Automated bone mapping			
features			
g. Ability to scan lumber spine (AP			
and Lateral), femur and forearm			
h. Supine lateral imaging			
i. Supine lateral lumbar spine			
densitometry for volumetric			
calculation of Bone Mineral Density			
(BMD)			
j. Capable of performing whole			
body scans			
k. On/Off positions shall be clearly			
identified or has indicator light			
l. Visible indication to identify that			·
it is ready to do exposure	<u> </u>		
m. Radiation symbol or indicator to			
denote exposure			-
n. Warning Signal to indicate			
termination of the exposure			-
o. Electronic timer			

AGENCY SPECIFICATIONS	BIDDER'S STATEMENT OF COMPLIANCE	ACTUAL OFFER	REFERENCE
p. Display of kV and mA			
q. Position indicator: laser light			
r. Spine phantom for QA/ QC			
s. Standard Software			
t. Basic Skeletal Package			
t.1) AP Spine			
t.2) Femur			
t.3) Dual Femur			
t.4) Forearm			
t.5) FRAX Fracture Risk Tool			
u. Total Body BMD			
v. Pediatric Package			
v.1) Pediatric AP Spine			i
v.2) Pediatric Total Body			
v.3) Pediatric Femur			
v.4) Pediatric Total Body – Birth to 20 years			
w. Orthopedic			
w.1) Orthopedic Hip			
w.2) Orthopedic Knee			
x. Other Software			
x.1) Digital Vertebral Assessment			
x.2) Advanced Hip Assessment			
x.3) Spine Geometry			
x.4) Hand, Encore			
x.5) Total Body Composition			
x.6) Advance Body Composition			
y. Connectivity: HL7, DICOM, Multi User DB (1-3)			
z. Work flow: Tele densitometry, Scan check, Report Composer			
aa. Accessories:	-		
aa.1) Block Phantoms for spine and whole body			

AGENCY SPECIFICATIONS	BIDDER'S STATEMENT OF COMPLIANCE	ACTUAL OFFER	REFERENCE
aa.2) Complete Table pad and			
positioning accessories			
aa.3) Desk top computer with at			
least 20 inch monitor and with Pre-			
installed latest Operating System, 64			
bit OEM, 4-physical cores, 4 GB Memory, 1 TB Hard Disc, DVD+/-RW			
Sata Drive, Tower Case with power			
supply unit 600W Max, USB			
keyboard and Mouse, USP 220-			
240V, At least 1 KVA, Productivity			
Software, Internet Security			
aa.4) One (1) unit Mobile computer			
table/cart			
aa.5) Printer: Color Inkjet			·
Resolution: 1200 x 1200 dpi	i		
Power Specifications: 110 volts			
C. ADDITIONAL REQUIREMENTS			
I. WARRANTY			
a. Comprehensive Warranty			
Certificate for (1) year on parts and			
three (3) years on service with			
Service Level Agreement (SLA) after			
testing and acceptance registration			
to the Philippine Nuclear Research			
Institute (PNRI) and Department of Health- Food and Drug		•	
Administration – Center for Device			·
Regulation Radiation Health and			
Research (DOH-FDA-CDRRHR).		·	
Reckoning of the warranty period			
will be upon approval of DOH-FDA-	•		
CDRRHR.			
b. The Service Level Agreement (SLA)			
shall cover the complete unit/system			
its sub-systems, components,			
associated accessories and			
peripherals supplied by third party			
should be considered by the bidder			
as its own. Warranty shall be signed by the manufacturer and must			
provide the guarantee that failures			
in materials and workmanship that			
The trials and working that	1		

	BIDDER'S		د
AGENCY SPECIFICATIONS	STATEMENT OF COMPLIANCE	ACTUAL OFFER	REFERENCE
occur within the warranty period will	COIVII LIAITEL		
be corrected. Such failures will			
include those attributable to			
abnormal aging. The maintenance	,		
and service of third party items will	•		
also be the sole responsibility of the			
primary vendor. Essential non-			
propriety spare parts should be			
made available.			
The SLA should cover the following:	•		
a. Guaranteed up-time of at least 95%			:
b. Availability of One (1) Service			·
Engineer assigned within Metro Manila			
c. Response Time: Within twenty			
four (24) hours from notice	!		
d. Mode of Delivery of service- with			
help desk that can be contacted by			
email, text and phone; and remote			
online troubleshooting.			
c. Supplier must specify post	-		
warranty comprehensive preventive			
maintenance costs including list and			
prices of major spare parts of the			
SPECT CT Scan and DEXA Bone			
Denstitometer and all accessories for			
the next three (3) years after the warranty period.			
II. TRAINING			
a. Two (2) weeks on-site training of			
three (3) Nuclear Medicine			
Technologists			
b. Two (2) weeks on-site training of			
one (1) Nuclear Medicine			
Physician			
III. DELIVERY PERIOD			
One hundred eighty (180) Calendar			
Days from receipt of Notice to			
Proceed including the Delivery,			
Installation, Testing, and			
Commissioning. Partial Delivery			
allowed within the completion/			
delivery period.	<u> </u>		

AGENCY SPECIFICATIONS	BIDDER'S STATEMENT OF COMPLIANCE	ACTUAL OFFER	REFERENCE
IV. DRAWING			
Please refer to the Drawing attached			
as Annex "I".			

I hereby certify that the statement of compliance to the foregoing technical specifications are true and correct, otherwise, if found to be false either during bid evaluation or post-qualification, the same shall give rise to automatic disqualification of our bids.

Name of Company	Signature Over Printed Name of	Date
	Authorized Representative	