REFERENCE OF TENDER	DESCRIPTION OF TENDER	TIME PERIOD OF TENDER	DEPARTMENT/ DIVISION/UNIT REQUESTING TENDER	FEES	CLOSING DATE NOT LATER THAN 2.00PM	FOCAL PERSON
KK/139/2025/HTD	PROVISION OF ULTRASOUND SYSTEMS FOR RADIOLOGY DEPARTMENT, MINISTRY OF HEALTH		DEPARTMENT OF HEALTHCARE TECHNOLOGY	\$500.00	24 ^{⊤H} June 2025	Muhd Amirul Fazleen bin Haji Khalidin Biomedical Engineer Suri Seri Begawan Hospital, Kuala Belait Ministry of Health Negara Brunei Darussalam Contact No: 3335331 ext 4222 email : fazleen.khalidin@moh.gov.bn

TENDER REFERENCE NO.:KK/139/2025/HTD

MINISTRY OF HEALTH NEGARA BRUNEI DARUSSALAM

PROVISION OF ULTRASOUND SYSTEMS FOR RADIOLOGY DEPARTMENT, MINISTRY OF HEALTH

- **TENDER FEES** : \$500.00
- RECEIPT NO. :
- CLOSING DATE : ON TUESDAY, 24TH JUNE 2025
- TIME : 2.00 PM
- FOA :

THE CHAIRMAN MINI TENDER BOARD, TENDER BOX GROUND FLOOR, MINISTRY OF HEALTH COMMONWEALTH DRIVE BANDAR SERI BEGAWAN BB3910 NEGARA BRUNEI DARUSSALAM

(NON CLUSTERING)

SPECIFICATIONS AND REQUIREMENTS

TENDER REFERENCE NO: KK/139/2025/HTD

NO.	TERMS AND CONDITIONS
1	Tenderer must be registered with the Ministry of Health.
2	TENDER FORM should be filled completely including the USER REQUIREMENT FORM (if available). Submission of incomplete form <u>MAY</u> cause DISQUALIFICATION OF TENDER .
3	Each tenderer is allowed to quote ONE BRAND/ MODEL WITH ONE PRICE ONLY for each item. Submission of more than one brand/model and price will cause DISQUALIFICATION OF TENDER.
4	Tenderers are required to submit individual proposal booklets for each item listed . Each item shall be treated as a standalone submission
5	All consumables supplied throughout this tender <u>shall</u> have a minimum expiry date of twelve (12) months / on delivery (if applicable). Should the consumables be urgently needed, provision of consumables with expiry date of less than twelve (12) months should be first agreed by the User before delivery is made (if applicable).
6	Brochures / catalogues should be submitted / attached with tender document.
7	Any room renovation which may be required, it is mandatory to conduct site visit (if applicable)
8	Samples should be submitted together with tender or within fourteen (14 days) of the tender closing dates (if applicable).
9	DELIVERY PERIOD: (Please state) Not More Than 90 days upon confirmation
10	PRICE VALIDITY: The quotation shall remain valid for 12 MONTHS from the final date for the submission of the quotation and no supplier may withdraw his/her quotation within that period. The Government reserves the right to extend this period if deemed necessary provided that such extension to the quotation validity period shall have written consent of the supplier(s).
11	The equipment supplied must be newly manufactured , unused, and in its original, sealed packaging. The equipment must not be previously owned, refurbished, or reconditioned in any form.
12	The vendor is required to provide proof of manufacture date and official certification from the original manufacturer confirming the equipment is new .
13	To provide justification for the price increase of a product previously supplied to the Ministry of Health by the same supplier/distributor

SCOPE OF WORK AND SUMMARY OF PRICES

This tender is for the supply of Ultrasound machines under a non-clustering approach for the following items:

Description	Quantity
Item 1: High-End Ultrasound System	3 Units
Item 2: Breast Imaging Ultrasound System	2 Units
Item 3: Mid-End Ultrasound System	1 Unit
Item 4: O&G Ultrasound System	3 Units
Item 5: Ultrasound Examination Couch and Chair	6 Units

ITEM 1	HIGH-END ULTRASOUND SYSTEM
1	SYSTEM ARHITECTURE
REF. NO.	DESCRIPTION
1	Three (3) units of high-end, cart-based ultrasound (US) system.
2	High-end ultrasound system with advanced imaging features such as high-resolution imaging, needle enhancement, elastography, AI-assisted imaging analysis. Support for contrast agents with specific pre-sets and quantification tools.
3	Advanced clinical applications: For comprehensive radiology studies for both adult and paediatric patients including abdominal, vascular, obstetrics, gynaecology, small parts, transcranial, paediatric and neonatal, musculoskeletal, urological and cardiac applications.
4	System must be able to provide exceptional clarity, resolution and depth of penetration.
5	The system is fully digital, compact, lightweight and easy to move.
6	Comes with independent steer and central locking wheels.
7	Comes with front and rear handles.
8	Has at least four (4) active transducer ports and one (1) parking probe port.
9	Has alphanumeric soft keys physical keyboard with easy access scans controls and facility to sanitise the system's keyboard to avoid cross contamination.
10	Has trackball for moving cursor
11	Inclusive of two (2) gel bottle holders.
12	Inclusive of one (1) unit of integrated gel warmer.
13	Has integrated cable management.
14	System storage: integrated SSD or HDD or both.
15	System storage: at least 1.5TB in total
16	Operating system: Window 10 or latest
17	Operating system: install on SSD
18	Has cybersecurity program to protect system
19	Power: able to operate on AC and on battery for at least 60 minutes or better
20	Battery: Self charging and has low-battery indicator
21	External data interface such as integrated DVD-RW drive, USB or its equivalent to enable external images transfer manually.
22	Capable to compare images from previous study.
23	Multiple preloaded application pre-sets available.
24	Control Monitor/Touch Panel Type: Touchscreen colour display.

ITEM 1	HIGH-END ULTRASOUND SYSTEM
1	SYSTEM ARHITECTURE
REF. NO.	DESCRIPTION
25	Control Monitor/Touch Panel Size: at least 9 inches or better
26	Control Monitor/Touch Panel has back-lighting
27	Display Viewing Monitor Type: LCD technology or better
28	Display Viewing Monitor Size: at least 22" or better
29	Display Viewing Monitor Resolution: Full HD (1920 x 1080) or better
30	Display Viewing Monitor has adjustable arm: can adjust tilt, swivel and height.
31	Display Viewing Monitor has adjustable contrast and brightness
32	Display Viewing Monitor can be fold down and locked for transportation.
33	Labelling and markings shall be clear legible and durable. (Durable to withstand routine cleaning)
34	OPERATING MODES The proposed system must include but not limited to the following operating modes:
34.1	Brightness-mode (B) for fundamental imaging mode for anatomical visualization.
34.2	Motion-mode (M) for motion analysis.
34.3	Trapezoidal mode for extended field of view for linear probes.
34.4	Pulsed wave (PW) mode for assessment of blood flow velocities
34.5	Continuous wave (CW) doppler mode for analysing high-velocity blood flow
34.6	Colour flow (CF) mode for mapping blood flow direction and velocity.
34.7	Power doppler imaging (PDI) mode for detecting low-velocity blood flow.
34.8	Contrast and tissue harmonic imaging for enhancing imaging for vascular studies and for clearer visualization of structures.
34.9	Spatial compounding for improving image quality by combining multiple beam angles.
34.10	Extended and convex field of view for enhancing visualization for larger areas during scanning
34.10	Speckle reduction imaging for reducing noises for clearer images
34.11	Automatic image optimisation for B-mode and doppler.
34.12	2D shear wave elastography for quantitative assessment of liver stiffness for fibrosis and Ultrasound-Derived Fat Fraction (UDFF) feature to measure the amount of fat within the liver tissues.
34.13	Adjustment of focal zone, gain, TGC, dynamic range, edge enhancement, gamma correction.
34.14	Frequency tuning for probes to adjust the frequency range on all transducers.

ITEM 1	HIGH-END ULTRASOUND SYSTEM
1	SYSTEM ARHITECTURE
REF. NO.	DESCRIPTION
34.15	Operator controlled multiple and adjustable focal zones.
34.16	Real-time pan zoom capability
34.17	Tools for labelling and anatomical markers for image annotation
34.18	Customizable user defined pre-sets for variety of clinical applications.
34.19	2D cine memory and M-mode scroll memory.
34.20	Cine loop facility up to one minute at 25 frames per second (fps).
34.21	Fast random-access image review.
34.22	Ability to apply wide variety of image processing after the examination.
34.23	Full size and split-screen display.
34.24	Support the following simultaneous display capability: B/PW
34.25	Support the following simultaneous display capability: B/CF
34.26	Support the following simultaneous display capability: B/spatial compounding
34.27	Support the following simultaneous display capability: B/PDI
34.28	Support the following simultaneous display capability: B/M
34.30	Multi-image split-screen display capability with combination such as: Live and frozen image comparisons
34.31	Multi-image split-screen display capability with combination such as: B or spatial compounding and B or spatial compounding
34.32	Multi-image split-screen display capability with combination such as: B or spatial compounding and CF
34.33	Multi-image split-screen display capability with combination such as: B or spatial compounding and PDI
34.34	Multi-image split-screen display capability with combination such as: PW/M
35	MEASUREMENT AND CALCULATION PACKAGES The proposed system shall include the following:
35.1	Basic measurements tools such as: distance (four points), angle (two points), calculation of diameter, area, circumference, calculation of volume by height/width/length, planimetry, automatic planimetry and empirical method.
35.2	Automated measurement packages including reports for abdominal, urology, renal, cardiac (M-mode and B-mode), obstetrics and gynaecological, vascular and blood flow, tissue elastography, musculoskeletal measurements, 3D/4D volume.
35.3	Real-time automatic doppler calculations and waveform indices.
35.4	Tools for hip dysplasia calculations including angle and joint measurements.

ITEM 1	HIGH-END ULTRASOUND SYSTEM
1	SYSTEM ARHITECTURE
REF. NO.	DESCRIPTION
35.5	Shear wave elastography measurement of liver stiffness in both m/s and kPa with automated interquartile range (IQR) and median value.
35.6	Inclusive of automatic foetal biometry tools measuring key foetal parameters like head circumference (HC), abdominal circumference (AC), femur length (FL) and biparietal diameter (BPD).
35.7	Inclusive of automated carotid intima-media thickness (CIMT) measurement tools.
35.8	Support panoramic imaging up to 60cm and curved imaging up to 360°
35.9	Inclusive of Shear wave elastography (SWE) and advanced strain imaging features for both quantitative and qualitative elastography and to measure tissue deformation in response to compression, ideal for small parts like breast and thyroid.
35.10	Inclusive of quantitative assessment of liver fat content using ultrasound or ultrasound-derived fat fraction (UDFF).
36	NETWORKING, CONNECTIVITY AND PACS INTEGRATION
36.1	Must fully support and comply to DICOM 3.0 standards, including: 1. DICOM Image Storage, 2. DICOM Modality Worklist (MWL), 3. DICOM Query/Retrieve (Q/R) and 4. DICOM Structured Reporting (SR)
36.2	Must support Modality Performed Procedure Step (MPPS)
36.4	Allow bidirectional data exchange for seamless workflow.
36.5	Image must be able to store as DICOM and PC format
36.6	System must have wireless connectivity and ethernet RJ45
36.7	System must be HL7 ready and activated
36.8	The systems proposed shall be integrated and connected to Radiology PACS (RPACS) therefore ALL licenses for integration with PACS should be included.
	Tenderer to include the cost of PACS integration quoted from PACS vendor.
	Please refer to PACS vendor for information on PACS integration requirement. Tenderer may contact the quotation of PACS integration for this project directly to the MOH's PACS vendor or through the Healthcare Technology Department, Ministry of Health

2	TRANSDUCERS
REF. NO.	DESCRIPTION
	The following transducer shall be included for each of the three high end ultrasound system:
1	Three (3) units of high frequency linear transducer for vascular, small parts, breast and musculoskeletal. (Tenderer to state the frequency.)
2	Three (3) units of broadband curved array transducer for general purpose abdominal and obstetric. (Tenderer to state the frequency.)
3	Three (3) units of broadband phased array transducer (micro-convex or sector) for cardiac echo and paediatric transcranial ultrasound. (Tenderer to state the frequency.)
4	Three (3) units of broadband curved array transducer for liver shear wave elastography (applicable only if a separate probe from the above is required for this purpose)
5	Three (3) units of linear array hockey stick transducer for peripheral vascular, small parts and superficial structures. (Tenderer to state the frequency.)
6	One (1) unit of curved intra-cavitary transducer for transvaginal obstetrics and gynaecology scan for RIPASH Radiology. Tenderer to state the frequency.
7	All transducers supplied must have a warranty period of at least one (1) year or better and comes with the necessary probe cover (if applicable)

3	DELIVERY AND INSTALLATION SITES
REF. NO.	DESCRIPTION
	Distribution for delivery, installation, testing and commissioning sites are as follows:
1	One (1) unit for Main ultrasound room, Radiology Department, Raja Isteri Pengiran Anak Saleha (RIPAS) Hospital, Brunei Muara.
2	One (1) unit for Radiology Unit, Pengiran Muda Mahkota Pengiran Muda Haji Al-Muhtadee Billah (PMMPMHAMB) Hospital, Tutong
3	One (1) unit for Radiology Unit, Suri Seri Begawan (SSB) Hospital, Kuala Belait

4	WARRANTY AND 3 YEARS POST WARRANTY PREVENTIVE MAINTENANCE (PM)
REF. NO.	DESCRIPTION
1	Tenderer to include warranty period of at least two (2) years
2	Tenderers to ACKNOWLEDGE the Warranty Undertaking Form in Section 4 stating the terms of warranty provided for the equipment in the tender for the period of two years. This includes but not limited to:
2.1	Scope of Warranty
2.2	Two time Planned Preventive Maintenance during warranty [Manufacturer's Standard]
3	Tenderer to provide additional 3 years Preventive Maintenance service post warranty [At least One Time PM per year or as per Manufacturer's Standard]
4	Tenderer to provide and attach the scope of work for 3 years PM service post warranty

5	END USER TRAINNING
REF. NO.	DESCRIPTION
1	Inclusive of clinical application training for all radiologists for at least seven (7) working days on-site for all the operation and applications offered with the system by Manufacturer's Application Specialist which includes but not limited to:
1.1	Basic user operation, including image transfers.
1.2	Application training for the use of the various applications provided with the system.
1.3	Basic maintenance, including troubleshooting
2	Training certificate must be provided by manufacturer or tenderer after completion of training sessions.
3	On-site follow up application training by application specialist after three (3) months of clinical use to ensure the system is fully optimised.
4	Two (2) sets of User/Operation Manual in English
5	Two (2) sets of Training Manual in English
6	Introductory Technical Training to Biomedical Engineers and Technicians at BME RIPASH Office by competent Tenderer's Engineer/Technicians that includes but not limited to:
6.1	Troubleshooting and basic corrective maintenance
6.2	Handling and basic inspection maintenance

6	INTERNATIONAL STANDARDS
REF. NO.	DESCRIPTION
	Equipment model offered must at least be in compliance and approved with four (4) of the following standards or better: (Please provide the documentation/certification to support this compliance)
1	European Union (CE MARK)
2	Food and Drug Administration (FDA) USA
3	International Electrotechnical Commission (IEC) 60601
4	International Electrotechnical Commission (IEC) 62304
5	International Organization for Standardization (ISO) 14971
6	International Organization for Standardization (ISO) 13485
7	International Organization for Standardization (ISO) 10993

7	DEMO UNIT
REF. NO.	DESCRIPTION
1	Tenderer to arrange a demo unit of the proposed system to be use for at least one to two (2) weeks – in Radiology Department, RIPASH.
	Please provide details of arranged demo unit

ITEM 2	BREAST IMAGING ULTRASOUND SYSTEM
1	SYSTEM ARHITECTURE
REF. NO.	DESCRIPTION
1	Two (2) units of cart-based ultrasound (US) system.
2	Configured specifically for breast and small parts imaging.
3	The system is fully digital, compact, lightweight and cart-based design
4	Comes with independent steer and central locking wheels.
5	Comes with front and rear handles.
6	Has at least three (3) active transducer ports and one (1) parking probe port.
7	Has alphanumeric soft keys physical keyboard with easy access scans controls and facility to sanitise the system's keyboard to avoid cross contamination.
8	Has trackball for moving cursor
9	Inclusive of two (2) gel bottle holders.
10	Inclusive of one (1) unit of integrated gel warmer.
11	Has integrated cable management.
12	System storage: integrated SSD or HDD or both.
13	System storage: at least 1.5TB in total
14	Operating system: Window 10 or latest
15	Operating system: install on SSD
16	Has cybersecurity program to protect system
17	Power: able to operate on AC and on battery for at least 60 minutes or better
18	Battery: Self charging and has low-battery indicator
19	External data interface such as integrated DVD-RW drive, USB or its equivalent to enable external images transfer manually.
20	Capable to compare images from previous study.
21	Multiple preloaded application pre-sets available.
22	Control Monitor/Touch Panel Type: Touchscreen colour display.
23	Control Monitor/Touch Panel Size: at least 9 inches or better
24	Control Monitor/Touch Panel has back-lighting
25	Display Viewing Monitor Type: LCD technology or better
26	Display Viewing Monitor Size: at least 22" or better

ITEM 2	BREAST IMAGING ULTRASOUND SYSTEM
1	SYSTEM ARHITECTURE
REF. NO.	DESCRIPTION
27	Display Viewing Monitor Resolution: Full HD (1920 x 1080) or better
28	Display Viewing Monitor has adjustable arm: can adjust tilt, swivel and height.
29	Display Viewing Monitor has adjustable contrast and brightness
30	Display Viewing Monitor can be fold down and locked for transportation.
31	Labelling and markings shall be clear legible and durable. (Durable to withstand routine cleaning)
32	Can be fold down and locked for transportation.
33	OPERATING MODES The proposed system must include but not limited to the following operating modes:
33.1	Brightness-mode (B) for fundamental imaging mode for anatomical visualization.
33.2	Motion-mode (M) for motion analysis.
33.3	Trapezoidal mode for extended field of view for linear probes.
33.4	Pulsed wave (PW) mode for assessment of blood flow velocities
33.5	Continuous wave (CW) doppler mode for analysing high-velocity blood flow
33.6	Colour flow (CF) mode for mapping blood flow direction and velocity.
33.7	Power doppler imaging (PDI) mode for detecting low-velocity blood flow.
33.8	Contrast and tissue harmonic imaging for enhancing imaging for vascular studies and for clearer visualization of structures.
33.9	Spatial compounding for improving image quality by combining multiple beam angles.
33.10	Extended and convex field of view for enhancing visualization for larger areas during scanning
33.11	Speckle reduction imaging for reducing noises for clearer images
33.12	Automatic image optimisation for B-mode and doppler.
33.13	Adjustment of focal zone, gain, TGC, dynamic range, edge enhancement, gamma correction.
33.14	Frequency tuning for probes to adjust the frequency range on all transducers.
33.15	Operator controlled multiple and adjustable focal zones.
33.16	Real-time pan zoom capability
33.17	Tools for labelling and anatomical markers for image annotation
33.18	Customizable user defined pre-sets for variety of clinical applications.
33.19	2D cine memory and M-mode scroll memory.

ITEM 2	BREAST IMAGING ULTRASOUND SYSTEM
1	SYSTEM ARHITECTURE
REF. NO.	DESCRIPTION
33.20	Cine loop facility up to one minute at 25 frames per second (fps).
33.21	Fast random-access image review.
33.22	Ability to apply wide variety of image processing after the examination.
33.23	Full size and split-screen display.
33.24	Support the following simultaneous display capability: B/PW
33.25	Support the following simultaneous display capability: B/CF
33.26	Support the following simultaneous display capability: B/spatial compounding
33.27	Support the following simultaneous display capability: B/PDI
33.28	Support the following simultaneous display capability: B/M
33.29	Multi-image split-screen display capability with combination such as: Live and frozen image comparisons
33.30	Multi-image split-screen display capability with combination such as: B or spatial compounding and B or spatial compounding
33.31	Multi-image split-screen display capability with combination such as: B or spatial compounding and CF
33.32	Multi-image split-screen display capability with combination such as: B or spatial compounding and PDI
33.33	Multi-image split-screen display capability with combination such as: PW/M
34	MEASUREMENT AND CALCULATION PACKAGES The proposed system shall include the following:
34.1	Basic measurements such as: distance (four points), angle (two points), calculation of diameter, area, circumference, calculation of volume by height/width/length, planimetry, automatic planimetry and empirical method.
34.2	Automated lesion analysis tools: auto lesion outline, volume calculation, BI-RADS Assistant.
34.3	Real-time automatic doppler and vascular calculations and waveform indices.
34.4	Elastography Measurements:
34.4.1	Shear wave elastography (SWE) measurement for breast stiffness in both m/s and kPa with automated interquartile range (IQR) and median value.
34.4.2	Strain elastography (SE) to evaluate stiffness (elasticity) of breast tissue measure tissue deformation in response to compression and provide a color-coded or greyscale map to represent the stiffness.
34.5	Biopsy and interventional measurement tools such as needle paths, visibility, distance- to-target measurement, localisation and marking.
34.6	Inclusive of breast productivity package or breast-specific lesion measurements such as aspect ratio and tumour-to-nipple distance along with tools for evaluating breast productivity.
34.7	Inclusive of automated carotid intima-media thickness (CIMT) measurement tools.
34.8	Support panoramic imaging up to 60cm and curved imaging up to 360°

ITEM 2	BREAST IMAGING ULTRASOUND SYSTEM
1	SYSTEM ARHITECTURE
REF. NO.	DESCRIPTION
35	NETWORKING, CONNECTIVITY AND PACS INTEGRATION
35.1	Must fully support and comply to DICOM 3.0 standards, including: 1. DICOM Image Storage, 2. DICOM Modality Worklist (MWL), 3. DICOM Query/Retrieve (Q/R) and 4. DICOM Structured Reporting (SR)
35.2	Must support Modality Performed Procedure Step (MPPS)
35.4	Allow bidirectional data exchange for seamless workflow.
35.5	Image must be able to store as DICOM and PC format
35.6	System must have wireless connectivity and ethernet RJ45
35.7	System must be HL7 ready and activated
	The systems proposed shall be integrated and connected to Radiology PACS (RPACS) therefore ALL licenses for integration with PACS should be included. Tenderer to include the cost of PACS integration quoted from PACS vendor.
35.8	Please refer to PACS vendor for information on PACS integration requirement. Tenderer may contact the quotation of PACS integration for this project directly to the MOH's PACS vendor or through the Healthcare Technology Department, Ministry of Health

2	TRANSDUCERS
REF. NO.	DESCRIPTION
	The following transducers shall be included for each of the two breast imaging ultrasound systems:
1	Two (2) units of high frequency linear transducer for vascular , small parts , breast and musculoskeletal . (Tenderer to state the frequency.)
2	Two (2) units of broadband curved array transducer for general purpose ultrasound imaging. (Tenderer to state the frequency.)
3	Two (2) units of linear transducer for breast applications. (Tenderer to state the frequency.)
4	All transducers supplied must have a warranty period of at least one (1) year or better and comes with the necessary probe cover (if applicable)

3	DELIVERY AND INSTALLATION SITES
REF. NO.	DESCRIPTION
	Distribution for delivery, installation, testing and commissioning sites are as follows:
1	Two (2) units to Breast Imaging Centre, Raja Isteri Pengiran Anak Saleha (RIPAS) Hospital, Brunei Muara.

4	WARRANTY AND 3 YEARS POST WARRANTY PREVENTIVE MAINTENANCE (PM)
REF. NO.	DESCRIPTION
1	Tenderer to include warranty period of at least two (2) years
2	Tenderers to ACKNOWLEDGE the Warranty Undertaking Form in Section 4 stating the terms of warranty provided for the equipment in the tender for the period of two years. This includes but not limited to:
2.1	Scope of Warranty
2.2	Two time Planned Preventive Maintenance during warranty [Manufacturer's Standard]
3	Tenderer to provide additional 3 years Preventive Maintenance service post warranty [At least One Time PM per year or as per Manufacturer's Standard]
4	Tenderer to provide and attach the scope of work for 3 years PM service post warranty

5	END USER TRAINNING
REF. NO.	DESCRIPTION
1	Inclusive of clinical application training for all radiologists for at least seven (7) working days on-site for all the operation and applications offered with the system by Manufacturer's Application Specialist which includes but not limited to:
1.1	Basic user operation, including image transfers.
1.2	Application training for the use of the various applications provided with the system.
1.3	Basic maintenance, including troubleshooting
2	Training certificate must be provided by manufacturer or tenderer after completion of training sessions.
3	On-site follow up application training by application specialist after three (3) months of clinical use to ensure the system is fully optimised.
4	Two (2) sets of User/Operation Manual in English
5	Two (2) sets of Training Manual in English
6	Introductory Technical Training to Biomedical Engineers and Technicians at BME RIPASH Office by competent Tenderer's Engineer/Technicians that includes but not limited to:
6.1	Troubleshooting and basic corrective maintenance
6.2	Handling and basic inspection maintenance

6	INTERNATIONAL STANDARDS
REF. NO.	DESCRIPTION
	Equipment model offered must at least be in compliance and approved with four (4) of the following standards or better: (Please provide the documentation/certification to support this compliance)
1	European Union (CE MARK)
2	Food and Drug Administration (FDA) USA
3	International Electrotechnical Commission (IEC) 60601
4	International Electrotechnical Commission (IEC) 62304
5	International Organization for Standardization (ISO) 14971
6	International Organization for Standardization (ISO) 13485
7	International Organization for Standardization (ISO) 10993

7	DEMO UNIT
REF. NO.	DESCRIPTION
1	Tenderer to arrange a demo unit of the proposed system to be use for at least one to two (2) weeks – in Radiology Department, RIPASH.

ITEM 3	MID-END ULTRASOUND SYSTEM
1	SYSTEM ARHITECTURE
REF. NO.	DESCRIPTION
1	One (1) unit of cart-based ultrasound (US) system.
2	Configured specifically for angiography imaging
3	The system is fully digital, compact, lightweight and easy to move.
4	Comes with independent steer and central locking wheels.
5	Comes with front and rear handles.
6	Has at least two (2) active transducer ports and one (1) parking probe port.
7	Has alphanumeric soft keys physical keyboard with easy access scans controls and facility to sanitise the system's keyboard to avoid cross contamination.
8	Has trackball for moving cursor
9	Inclusive of two (2) gel bottle holders.
10	Inclusive of one (1) unit of integrated gel warmer.
11	Has integrated cable management.
12	System storage: integrated SSD or HDD or both.
13	System storage: at least 1.5TB in total
14	Operating system: Window 10 or latest
15	Operating system: install on SSD
16	Has cybersecurity program to protect system
17	Power: able to operate on AC and on battery for at least 60 minutes or better
18	Battery: Self charging and has low-battery indicator
19	External data interface such as integrated DVD-RW drive, USB or its equivalent to enable external images transfer manually.
20	Capable to compare images from previous study.
21	Multiple preloaded application pre-sets available.
22	Control Monitor/Touch Panel Type: Touchscreen colour display.
23	Control Monitor/Touch Panel Size: at least 9 inches or better
24	Control Monitor/Touch Panel has back-lighting
25	Display Viewing Monitor Type: LCD technology or better
26	Display Viewing Monitor Size: at least 21" or better

ITEM 3	MID-END ULTRASOUND SYSTEM
1	SYSTEM ARHITECTURE
REF. NO.	DESCRIPTION
27	Display Viewing Monitor Resolution: Full HD (1920 x 1080) or better
28	Display Viewing Monitor has adjustable arm: can adjust tilt, swivel and height.
29	Display Viewing Monitor has adjustable contrast and brightness
30	Display Viewing Monitor can be fold down and locked for transportation.
31	Labelling and markings shall be clear legible and durable. (Durable to withstand routine cleaning)
32	Can be fold down and locked for transportation.
33	OPERATING MODES The proposed system must include but not limited to the following operating modes:
33.1	Brightness-mode (B) for fundamental imaging mode for anatomical visualization.
33.2	Motion-mode (M) for motion analysis.
33.3	Trapezoidal mode for extended field of view for linear probes.
33.4	Pulsed wave (PW) mode for assessment of blood flow velocities
33.5	Continuous wave (CW) doppler mode for analysing high-velocity blood flow
33.6	Colour flow (CF) mode for mapping blood flow direction and velocity.
33.7	Power doppler imaging (PDI) mode for detecting low-velocity blood flow.
33.8	Contrast and tissue harmonic imaging for enhancing imaging for vascular studies and for clearer visualization of structures.
33.9	Spatial compounding for improving image quality by combining multiple beam angles.
33.10	Extended and convex field of view for enhancing visualization for larger areas during scanning
33.11	Speckle reduction imaging for reducing noises for clearer images
33.12	Automatic image optimisation for B-mode and doppler.
33.14	Adjustment of focal zone, gain, TGC, dynamic range, edge enhancement, gamma correction.
33.15	Frequency tuning for probes to adjust the frequency range on all transducers.
33.16	Operator controlled multiple and adjustable focal zones.
33.17	Real-time pan zoom capability
33.18	Tools for labelling and anatomical markers for image annotation
33.19	Customizable user defined pre-sets for variety of clinical applications.

ITEM 3	MID-END ULTRASOUND SYSTEM
1	SYSTEM ARHITECTURE
REF. NO.	DESCRIPTION
33.20	2D cine memory and M-mode scroll memory.
33.21	Cine loop facility up to one minute at 25 frames per second (fps).
33.22	Fast random-access image review.
33.23	Ability to apply wide variety of image processing after the examination.
33.24	Full size and split-screen display.
33.25	Support the following simultaneous display capability: B/PW
33.26	Support the following simultaneous display capability: B/CF
33.27	Support the following simultaneous display capability: B/spatial compounding
33.28	Support the following simultaneous display capability: B/PDI
33.29	Support the following simultaneous display capability: B/M
33.30	Multi-image split-screen display capability with combination such as: Live and frozen image comparisons
33.31	Multi-image split-screen display capability with combination such as: B or spatial compounding and B or spatial compounding
33.32	Multi-image split-screen display capability with combination such as: B or spatial compounding and CF
33.33	Multi-image split-screen display capability with combination such as: B or spatial compounding and PDI
33.34	Multi-image split-screen display capability with combination such as: PW/M
34	MEASUREMENT AND CALCULATION PACKAGES The proposed system shall include the following:
34.1	Basic measurements such as: distance (four points), angle (two points), calculation of diameter, area, circumference, calculation of volume by height/width/length, planimetry, automatic planimetry and empirical method.
34.2	Automated measurement packages including reports, urology, renal, cardiac (M-mode and B-mode), obstetrics and gynaecological and vascular.
34.3	Real-time automatic doppler calculations and waveform indices.
34.4	Support panoramic imaging up to 60cm and curved imaging up to 360 degrees
35	NETWORKING, CONNECTIVITY AND PACS INTEGRATION
35.1	Must fully support and comply to DICOM 3.0 standards, including: 1. DICOM Image Storage, 2. DICOM Modality Worklist (MWL), 3. DICOM Query/Retrieve (Q/R) and 4. DICOM Structured Reporting (SR)
35.2	Must support Modality Performed Procedure Step (MPPS)
35.3	Allow bidirectional data exchange for seamless workflow.

ITEM 3	MID-END ULTRASOUND SYSTEM
1	SYSTEM ARHITECTURE
REF. NO.	DESCRIPTION
35.4	Image must be able to store as DICOM and PC format
35.5	System must have wireless connectivity and ethernet RJ45
35.6	System must be HL7 ready and activated
36.7	The systems proposed shall be integrated and connected to Radiology PACS (RPACS) therefore ALL licenses for integration with PACS should be included.
	Tenderer to include the cost of PACS integration quoted from PACS vendor.
	Please refer to PACS vendor for information on PACS integration requirement. Tenderer may contact the quotation of PACS integration for this project directly to the MOH's PACS vendor or through the Healthcare Technology Department, Ministry of Health

2	TRANSDUCERS
REF. NO.	DESCRIPTION
	The following transducers shall be included for the mid-end ultrasound system:
1	One (1) units of high frequency linear transducer for vascular, small parts, breast and musculoskeletal. (Tenderer to state the frequency.)
2	One (1) units of broadband curved array transducer for general purpose abdominal and obstetric. (Tenderer to state the frequency.)
3	All transducers supplied must have a warranty period of at least one (1) year or better and comes with the necessary probe cover (if applicable)

3	DELIVERY AND INSTALLATION SITES
REF. NO.	DESCRIPTION
	Distribution for delivery, installation, testing and commissioning sites are as follows:
1	One (1) unit for Angiography Room, Radiology Department, Raja Isteri Pengiran Anak Saleha (RIPAS) Hospital, Brunei Muara.

4	WARRANTY AND 3 YEARS POST WARRANTY PREVENTIVE MAINTENANCE (PM)
REF. NO.	DESCRIPTION
1	Tenderer to include warranty period of at least two (2) years
2	Tenderers to ACKNOWLEDGE the Warranty Undertaking Form in Section 4 stating the terms of warranty provided for the equipment in the tender for the period of two years. This includes but not limited to:
2.1	Scope of Warranty
2.2	Two time Planned Preventive Maintenance during warranty [Manufacturer's Standard]
3	Tenderer to provide additional 3 years Preventive Maintenance service post warranty [At least One Time PM per year or as per Manufacturer's Standard]
4	Tenderer to provide and attach the scope of work for 3 years PM service post warranty

5	END USER TRAINNING
REF. NO.	DESCRIPTION
1	Inclusive of clinical application training for all radiologists for at least seven (7) working days on-site for all the operation and applications offered with the system by Manufacturer's Application Specialist which includes but not limited to:
1.1	Basic user operation, including image transfers.
1.2	Application training for the use of the various applications provided with the system.
1.3	Basic maintenance, including troubleshooting
2	Training certificate must be provided by manufacturer or tenderer after completion of training sessions.
3	On-site follow up application training by application specialist after three (3) months of clinical use to ensure the system is fully optimised.
4	Two (2) sets of User/Operation Manual in English

5	END USER TRAINNING
REF. NO.	DESCRIPTION
5	Two (2) sets of Training Manual in English
6	Introductory Technical Training to Biomedical Engineers and Technicians at BME RIPASH Office by competent Tenderer's Engineer/Technicians that includes but not limited to:
6.1	Troubleshooting and basic corrective maintenance
6.2	Handling and basic inspection maintenance

6	INTERNATIONAL STANDARDS
REF. NO.	DESCRIPTION
	Equipment model offered must at least be in compliance and approved with four (4) of the following standards or better: (Please provide the documentation/certification to support this compliance)
1	European Union (CE MARK)
2	Food and Drug Administration (FDA) USA
3	International Electrotechnical Commission (IEC) 60601
4	International Electrotechnical Commission (IEC) 62304
5	International Organization for Standardization (ISO) 14971
6	International Organization for Standardization (ISO) 13485
7	International Organization for Standardization (ISO) 10993

7	DEMO UNIT
REF. NO.	DESCRIPTION
1	Tenderer to arrange a demo unit of the proposed system to be use for at least one to two (2) weeks – in Radiology Department, RIPASH.

ITEM 4	O&G ULTRASOUND SYSTEM
1	SYSTEM ARHITECTURE
REF. NO.	DESCRIPTION
1	Real-time image optimization and transducer technology migration
2	Imaging frequency range: 1–20 MHz
3	At least 4 active transducer ports with electronic selection
4	Integrated gel warmer with temperature adjustment
5	Transducer storage holder for minimum 7 probes & utility storage baskets with gel holder
6	Integrated cable management with multiple cable hooks, front/back handles, and lockable wheels
7	Customizable imaging modes and parameters
8	IMAGING CAPABILITIES
8.1	Imaging modes: B-mode, M-mode, Color & Power Doppler, Pulsed Wave & Continuous Wave Doppler
8.2	Adjustable field of view size and position, 2D beam steering angle $\geq \pm 30^{\circ}$
8.3	Parenchymal image harmonization for reduced noise and improved sharpness
8.4	Speckle noise reduction with \geq 3 levels via dynamic tissue contrast enhancement
9	CONTROL PANEL & DISPLAY
9.1	Height-adjustable control panel with ±90° swivel articulation
9.2	Tiltable touchscreen interface with USB ports
9.2 9.3	Tiltable touchscreen interface with USB ports Conventional keyboard and laser optical trackball with backlighting
9.2 9.3 9.4	Tiltable touchscreen interface with USB ports Conventional keyboard and laser optical trackball with backlighting In-Plane Switching monitor
9.2 9.3 9.4 9.5	Tiltable touchscreen interface with USB ports Conventional keyboard and laser optical trackball with backlighting In-Plane Switching monitor Display size: ≥ 21 inches
9.2 9.3 9.4 9.5 9.6	Tiltable touchscreen interface with USB ports Conventional keyboard and laser optical trackball with backlighting In-Plane Switching monitor Display size: ≥ 21 inches Display resolution ≥ 1920×1080
9.2 9.3 9.4 9.5 9.6 9.7	Tiltable touchscreen interface with USB ports Conventional keyboard and laser optical trackball with backlighting In-Plane Switching monitor Display size: ≥ 21 inches Display resolution ≥ 1920×1080 Digital calipers: minimum 4 active
9.2 9.3 9.4 9.5 9.6 9.7 9.8	Tiltable touchscreen interface with USB portsConventional keyboard and laser optical trackball with backlightingIn-Plane Switching monitorDisplay size: ≥ 21 inchesDisplay resolution ≥ 1920×1080Digital calipers: minimum 4 activeZoom and pan on both real-time and frozen images
9.2 9.3 9.4 9.5 9.6 9.7 9.8 10	Tiltable touchscreen interface with USB ports Conventional keyboard and laser optical trackball with backlighting In-Plane Switching monitor Display size: ≥ 21 inches Display resolution ≥ 1920×1080 Digital calipers: minimum 4 active Zoom and pan on both real-time and frozen images POWER, SOFTWARE & STORAGE
9.2 9.3 9.4 9.5 9.6 9.7 9.8 10 10.1	Tiltable touchscreen interface with USB portsConventional keyboard and laser optical trackball with backlightingIn-Plane Switching monitorDisplay size: ≥ 21 inchesDisplay resolution ≥ 1920×1080Digital calipers: minimum 4 activeZoom and pan on both real-time and frozen imagesPOWER, SOFTWARE & STORAGEMains supply: 100–240V AC, 50/60 Hz
9.2 9.3 9.4 9.5 9.6 9.7 9.8 10 10.1 10.2	Tiltable touchscreen interface with USB ports Conventional keyboard and laser optical trackball with backlighting In-Plane Switching monitor Display size: ≥ 21 inches Display resolution ≥ 1920×1080 Digital calipers: minimum 4 active Zoom and pan on both real-time and frozen images POWER, SOFTWARE & STORAGE Mains supply: 100–240V AC, 50/60 Hz Battery integrated
9.2 9.3 9.4 9.5 9.6 9.7 9.8 10 10.1 10.2 10.3	Tiltable touchscreen interface with USB portsConventional keyboard and laser optical trackball with backlightingIn-Plane Switching monitorDisplay size: ≥ 21 inchesDisplay resolution ≥ 1920×1080Digital calipers: minimum 4 activeZoom and pan on both real-time and frozen imagesPOWER, SOFTWARE & STORAGEMains supply: 100–240V AC, 50/60 HzBattery integratedBattery operation: ≥ 70 minutes scan time with status display
9.2 9.3 9.4 9.5 9.6 9.7 9.8 10 10.1 10.2 10.3 10.4	Tiltable touchscreen interface with USB portsConventional keyboard and laser optical trackball with backlightingIn-Plane Switching monitorDisplay size: ≥ 21 inchesDisplay resolution ≥ 1920×1080Digital calipers: minimum 4 activeZoom and pan on both real-time and frozen imagesPOWER, SOFTWARE & STORAGEMains supply: 100–240∨ AC, 50/60 HzBattery integratedBattery operation: ≥ 70 minutes scan time with status displayOperating System: Windows 10 or later with antivirus installed

ITEM 4	O&G ULTRASOUND SYSTEM
1	SYSTEM ARHITECTURE
REF. NO.	DESCRIPTION
10.6	Cine memory: not less than 500 frames
11	NETWORKING, CONNECTIVITY AND PACS INTEGRATION
11.1	Must fully support and comply to DICOM 3.0 standards, including: 1. DICOM Image Storage, 2. DICOM Modality Worklist (MWL), 3. DICOM Query/Retrieve (Q/R) and 4. DICOM Structured Reporting (SR)
11.2	Must support Modality Performed Procedure Step (MPPS)
11.3	Allow bidirectional data exchange for seamless workflow.
11.4	Image must be able to store as DICOM and PC format
11.5	System must have wireless connectivity and ethernet RJ45
11.6	System must be HL7 ready and activated
11.7	The systems proposed shall be integrated and connected to Radiology PACS (RPACS) therefore ALL licenses for integration with PACS should be included. Tenderer to include the cost of PACS integration quoted from PACS vendor.
	Please refer to PACS vendor for information on PACS integration requirement. Tenderer may contact the quotation of PACS integration for this project directly to the MOH's PACS vendor or through the Healthcare Technology Department, Ministry of Health

1	TRANSDUCERS
REF. NO.	DESCRIPTION
	The following transducers shall be included for the O&G ultrasound system
1	One (1) unit of multi-frequency convex probe ■ Penetration ≥ 350 mm and Field of View ≥ 70 ⁰ (Tenderer to state the Model, Penetration, FOV and Frequency)
2	One (1) unit of Endo Cavity Transducer ■ penetration ≥ 140 mm and Field of View ≥ 220 ⁰ (Tenderer to state the Model, Penetration, FOV and Frequency)
3	One (1) unit of 4D Volume Transducer ■ penetration ≥ 300 mm and Field of View ≥ 65 ⁰ (Tenderer to state the Model, Penetration, FOV and Frequency)
4	One (1) unit of 4D Endo Cavity Volume Transducer ■ penetration ≥ 160 mm and Field of View ≥ 145 ⁰ (Tenderer to state the Model, Penetration, FOV and Frequency)
5	All transducers supplied must have a warranty period of at least one (1) year or better and comes with the necessary probe cover (if applicable)

2	DELIVERY AND INSTALLATION SITES	
REF. NO.	DESCRIPTION	
	Distribution for delivery, installation, testing and commissioning sites are as follows:	
1	One (1) unit for Pengkalan Batu Health Centre, MCH, Brunei Muara.	
2	One (1) unit for Sg Kelugos Health Centre, MCH, Tutong.	
3	One (1) unit for Telisai Health Centre, MCH, Tutong.	

3	WARRANTY AND 3 YEARS POST WARRANTY PREVENTIVE MAINTENANCE (PM		
REF. NO.	DESCRIPTION		
1	Tenderer to include warranty period of at least two (2) years		
2	Tenderers to ACKNOWLEDGE the Warranty Undertaking Form in Section 4 stating the terms of warranty provided for the equipment in the tender for the period of two years. This includes but not limited to:		
2.1	Scope of Warranty		
2.2	Two time Planned Preventive Maintenance during warranty [Manufacturer's Standard]		
3	Tenderer to provide additional 3 years Preventive Maintenance service post warranty [At least One Time PM per year or as per Manufacturer's Standard]		
4	Tenderer to provide and attach the scope of work for 3 years PM service post warranty		

3	END USER TRAINNING		
REF. NO.	DESCRIPTION		
1	Inclusive of clinical application training for all radiologists for at least seven (7) working days on-site for all the operation and applications offered with the system by Manufacturer's Application Specialist which includes but not limited to:		
1.1	Basic user operation, including image transfers.		
1.2	Application training for the use of the various applications provided with the system.		
1.3	Basic maintenance, including troubleshooting		
2	Training certificate must be provided by manufacturer or tenderer after completion of training sessions.		
3	On-site follow up application training by application specialist after three (3) months of clinical use to ensure the system is fully optimised.		
4	Two (2) sets of User/Operation Manual in English		
5	Two (2) sets of Training Manual in English		
6	Introductory Technical Training to Biomedical Engineers and Technicians at BME RIPASH Office by competent Tenderer's Engineer/Technicians that includes but not limited to:		
6.1	Troubleshooting and basic corrective maintenance		
6.2	Handling and basic inspection maintenance		

4	INTERNATIONAL STANDARDS		
REF. NO.	DESCRIPTION		
	Equipment model offered must at least be in compliance and approved with four (4) of the following standards or better: (Please provide the documentation/certification to support this compliance)		
1	European Union (CE MARK)		
2	Food and Drug Administration (FDA) USA		
3	International Electrotechnical Commission (IEC) 60601		
4	International Electrotechnical Commission (IEC) 62304		
5	International Organization for Standardization (ISO) 14971		
6	International Organization for Standardization (ISO) 13485		
7	International Organization for Standardization (ISO) 10993		

ITEM 5	ULTRASOUND EXAMINATION COUCH AND CHAIR		
1	SYSTEM ARHITECTURE		
REF. NO.	DESCRIPTION		
1	Six (6) units of ultrasound examination couch and Ultrasound operator chair		
2	Ultrasound Examination Couch: Electric height adjustable type		
3	Ultrasound Examination Couch: Adjustable back and leg rest.		
4	Ultrasound Examination Couch: Complete with disposable paper dispenser mounts to the head end of the couch.		
5	Ultrasound Examination Couch: Suitable dimension for dedicated room (Please refer to each location)		
6	Ultrasound Chair: Height adjustable ultrasound chair (saddle type) for the operators with wheels and adjustable back		

2	DELIVERY AND INSTALLATION SITES		
REF. NO.	DESCRIPTION		
	Distribution for delivery, installation, testing and commissioning sites are as follows:		
1	One (1) unit for Main ultrasound room, Radiology Department, Raja Isteri Pengiran Anak Saleha (RIPAS) Hospital, Brunei Muara.		
2	One (1) unit for Radiology Unit, Pengiran Muda Mahkota Pengiran Muda Haji Al- Muhtadee Billah (PMMPMHAMB) Hospital, Tutong		
3	One (1) unit for Radiology Unit, Suri Seri Begawan (SSB) Hospital, Kuala Belait		
4	One (1) unit for Angio Room, Radiology Department, Raja Isteri Pengiran Anak Saleha (RIPAS) Hospital, Brunei Muara.		
5	Two (2) units for Breast Imaging Centre, Raja Isteri Pengiran Anak Saleha (RIPAS) Hospital, Brunei Muara.		

3	WARRANTY
REF. NO.	DESCRIPTION
1	Tenderer to include warranty period of at least two (2) years
2	Tenderers to ACKNOWLEDGE the Warranty Undertaking Form in Section 4 stating the terms and scope of warranty provided for the equipment in the tender for the period of two years.

TENDER FORM

To:

TENDER REFERENCE NO: KK/139/2025/HTD

INVITATION TO TENDER PROVISION OF ULTRASOUND SYSTEMS FOR RADIOLOGY DEPARTMENT, MINISTRY OF HEALTH

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TENDER FORM

To:

TENDER REFERENCE NO: KK/139/2025/HTD

INVITATION TO TENDER PROVISION OF ULTRASOUND SYSTEMS FOR RADIOLOGY DEPARTMENT, MINISTRY OF HEALTH

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TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

TERMS AND CONDITIONS		
NO.	TERMS AND CONDITIONS	VENDOR'S OFFER
1	Tenderer must be registered with the Ministry of Health.	
2	TENDER FORM should be filled completely including the USER REQUIREMENT FORM (if available). Submission of incomplete form <u>MAY</u> cause DISQUALIFICATION OF TENDER .	
3	Each tenderer is allowed to quote ONE BRAND/ MODEL WITH ONE PRICE ONLY for each item. Submission of more than one brand/model and price will cause DISQUALIFICATION OF TENDER .	
4	Tenderers are required to submit individual proposal booklets for each item listed . Each item shall be treated as a standalone submission	
5	All consumables supplied throughout this tender <u>shall</u> have a minimum expiry date of twelve (12) months / on delivery (if applicable). Should the consumables be urgently needed, provision of consumables with expiry date of less than twelve (12) months should be first agreed by the User before delivery is made (if applicable).	
6	Brochures / catalogues should be submitted / attached with tender document.	
7	Any room renovation which may be required, it is mandatory to conduct site visit (if applicable)	
8	Samples should be submitted together with tender or within fourteen (14 days) of the tender closing dates (if applicable).	

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

TERMS AND CONDITIONS		
NO.	TERMS AND CONDITIONS	VENDOR'S OFFER
9	DELIVERY PERIOD: (Please state) Not More Than 90 days upon confirmation	
10	PRICE VALIDITY: The quotation shall remain valid for 12 MONTHS from the final date for the submission of the quotation and no supplier may withdraw his/her quotation within that period. The Government reserves the right to extend this period if deemed necessary provided that such extension to the quotation validity period shall have written consent of the supplier(s).	
11	The equipment supplied must be newly manufactured , unused, and in its original, sealed packaging. The equipment must not be previously owned, refurbished, or reconditioned in any form.	
12	The vendor is required to provide proof of manufacture date and official certification from the original manufacturer confirming the equipment is new .	
13	To provide justification for the price increase of a product previously supplied to the Ministry of Health by the same supplier/distributor	

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

SCOPE OF WORK AND SUMMARY OF PRICES						
This tender is for the supply of Ultrasound machines under a non- clustering approach for the following items:		VES	NO			
Description	Quantity	TES		UNIT PRICE		
Item 1: High-End Ultrasound System	3 Units			BND\$	BND\$	
Item 2: Breast Imaging Ultrasound System	2 Units			BND\$	BND\$	
Item 3: Mid-End Ultrasound System	1 Unit			BND\$	BND\$	
Item 4: O&G Ultrasound System	3 Units			BND\$	BND\$	
Item 5: Ultrasound Examination Couch and Chair	6 Units			BND\$	BND\$	

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

ITEM 1: HIGH-END ULTRASOUND SYSTEM						
SECTION 1 – USER REQUIREMENTS						
1	1 SYSTEM ARHITECTURE					
REF. NO.	DESCRIPTION		(√)	STATE OR SPECIFY OR REMARKS OR BROCHURE PAGE		
			NO			
1	Three (3) units of high-end, cart-based ultrasound (US) system.					
2	High-end ultrasound system with advanced imaging features such as high-resolution imaging, needle enhancement, elastography, AI-assisted imaging analysis. Support for contrast agents with specific pre-sets and quantification tools.					
3	Advanced clinical applications: For comprehensive radiology studies for both adult and paediatric patients including abdominal, vascular, obstetrics, gynaecology, small parts, transcranial, paediatric and neonatal, musculoskeletal, urological and cardiac applications.					
4	System must be able to provide exceptional clarity, resolution and depth of penetration.					
5	The system is fully digital, compact, lightweight and easy to move.					
6	Comes with independent steer and central locking wheels.					
7	Comes with front and rear handles.					
8	Has at least four (4) active transducer ports and one (1) parking probe port.					
9	Has alphanumeric soft keys physical keyboard with easy access scans controls and facility to sanitise the system's keyboard to avoid cross contamination.					
10	Has trackball for moving cursor					

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

ITEM 1: HIGH-END ULTRASOUND SYSTEM					
SECTION 1 – USER REQUIREMENTS					
1	1 SYSTEM ARHITECTURE				
REF. NO.	DESCRIPTION	Tick (√)		STATE OR SPECIFY OR	
		YES	NO	PAGE	
11	Inclusive of two (2) gel bottle holders.				
12	Inclusive of one (1) unit of integrated gel warmer.				
13	Has integrated cable management.				
14	System storage: integrated SSD or HDD or both.				
15	System storage: at least 1.5TB in total				
16	Operating system: Window 10 or latest				
17	Operating system: install on SSD				
18	Has cybersecurity program to protect system				
19	Power: able to operate on AC and on battery for at least 60 minutes or better				
20	Battery: Self charging and has low-battery indicator				
21	External data interface such as integrated DVD-RW drive, USB or its equivalent to enable external images transfer manually.				

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

ITEM 1: HIGH-END ULTRASOUND SYSTEM					
SECTION 1 – USER REQUIREMENTS					
1	SYSTEM ARHITECTURE				
REF. NO.	DESCRIPTION	Tick (✓)		STATE OR SPECIFY OR	
		YES	NO	PAGE	
22	Capable to compare images from previous study.				
23	Multiple preloaded application pre-sets available.				
24	Control Monitor/Touch Panel Type: Touchscreen colour display.				
25	Control Monitor/Touch Panel Size: at least 9 inches or better				
26	Control Monitor/Touch Panel has back-lighting				
27	Display Viewing Monitor Type: LCD technology or better				
28	Display Viewing Monitor Size: at least 22" or better				
29	Display Viewing Monitor Resolution: Full HD (1920 x 1080) or better				
30	Display Viewing Monitor has adjustable arm: can adjust tilt, swivel and height.				
31	Display Viewing Monitor has adjustable contrast and brightness				
32	Display Viewing Monitor can be fold down and locked for transportation.				
TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

ITEM 1: HIGH-END ULTRASOUND SYSTEM				
	SECTION 1 – USER REQUIREMENTS			
1	SYSTEM ARHITECTURE			
REF.	DESCRIPTION	Tick (√)		STATE OR SPECIFY OR
NO.	IO. YE	YES	NO	PAGE
33	Labelling and markings shall be clear legible and durable. (Durable to withstand routine cleaning)			
34	OPERATING MODES The proposed system must include but not limited to the following operating modes:			
34.1	Brightness-mode (B) for fundamental imaging mode for anatomical visualization.			
34.2	Motion-mode (M) for motion analysis.			
34.3	Trapezoidal mode for extended field of view for linear probes.			
34.4	Pulsed wave (PW) mode for assessment of blood flow velocities			
34.5	Continuous wave (CW) doppler mode for analysing high-velocity blood flow			
34.6	Colour flow (CF) mode for mapping blood flow direction and velocity.			
34.7	Power doppler imaging (PDI) mode for detecting low-velocity blood flow.			
34.8	Contrast and tissue harmonic imaging for enhancing imaging for vascular studies and for clearer visualization of structures.			
34.9	Spatial compounding for improving image quality by combining multiple beam angles.			

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

ITEM 1: HIGH-END ULTRASOUND SYSTEM						
	SECTION 1 – USER REQUIREMENTS					
1	SYSTEM ARHITECTURE					
REF.	DESCRIPTION	Tick	(•)	STATE OR SPECIFY OR		
NO.	DESCRIPTION	YES	NO	PAGE		
34.10	Extended and convex field of view for enhancing visualization for larger areas during scanning					
34.10	Speckle reduction imaging for reducing noises for clearer images					
34.11	Automatic image optimisation for B-mode and doppler.					
34.12	2D shear wave elastography for quantitative assessment of liver stiffness for fibrosis and Ultrasound-Derived Fat Fraction (UDFF) feature to measure the amount of fat within the liver tissues.					
34.13	Adjustment of focal zone, gain, TGC, dynamic range, edge enhancement, gamma correction.					
34.14	Frequency tuning for probes to adjust the frequency range on all transducers.					
34.15	Operator controlled multiple and adjustable focal zones.					
34.16	Real-time pan zoom capability					
34.17	Tools for labelling and anatomical markers for image annotation					
34.18	Customizable user defined pre-sets for variety of clinical applications.					
34.19	2D cine memory and M-mode scroll memory.					

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

	ITEM 1: HIGH-END ULTRASOUND SYSTEM				
	SECTION 1 – USER REQUIREMENTS				
1	SYSTEM ARHITECTURE				
REF.	DECODIDITION	Tick (√)		STATE OR SPECIFY OR	
NO.	DESCRIPTION	YES	NO	PAGE	
34.20	Cine loop facility up to one minute at 25 frames per second (fps).				
34.21	Fast random-access image review.				
34.22	Ability to apply wide variety of image processing after the examination.				
34.23	Full size and split-screen display.				
34.24	Support the following simultaneous display capability: B/PW				
34.25	Support the following simultaneous display capability: B/CF				
34.26	Support the following simultaneous display capability: B/spatial compounding				
34.27	Support the following simultaneous display capability: B/PDI				
34.28	Support the following simultaneous display capability: B/M				
34.30	Multi-image split-screen display capability with combination such as: Live and frozen image comparisons				
34.31	Multi-image split-screen display capability with combination such as: B or spatial compounding and B or spatial compounding				

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

ITEM 1: HIGH-END ULTRASOUND SYSTEM						
	SECTION 1 – USER REQUIREMENTS					
1	1 SYSTEM ARHITECTURE					
REF.	DESCRIPTION	Tick	(✓)	STATE OR SPECIFY OR		
NO.	DESCRIPTION	YES	NO	REMARKS OR BROCHURE O PAGE		
34.32	Multi-image split-screen display capability with combination such as: B or spatial compounding and CF					
34.33	Multi-image split-screen display capability with combination such as: B or spatial compounding and PDI					
34.34	Multi-image split-screen display capability with combination such as: PW/M					
35	MEASUREMENT AND CALCULATION PACKAGES The proposed system shall include the following:					
35.1	Basic measurements tools such as: distance (four points), angle (two points), calculation of diameter, area, circumference, calculation of volume by height/width/length, planimetry, automatic planimetry and empirical method.					
35.2	Automated measurement packages including reports for abdominal, urology, renal, cardiac (M-mode and B-mode), obstetrics and gynaecological, vascular and blood flow, tissue elastography, musculoskeletal measurements, 3D/4D volume.					
35.3	Real-time automatic doppler calculations and waveform indices.					
35.4	Tools for hip dysplasia calculations including angle and joint measurements.					
35.5	Shear wave elastography measurement of liver stiffness in both m/s and kPa with automated interquartile range (IQR) and median value.					

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

ITEM 1: HIGH-END ULTRASOUND SYSTEM						
	SECTION 1 – USER REQUIREMENTS					
1	SYSTEM ARHITECTURE					
REF.	DESCRIPTION	Tick	(√)	STATE OR SPECIFY OR		
NO.	DESCRIPTION	YES	NO	(✓) STATE OR SPECIFY OR REMARKS OR BROCHURE PAGE		
35.6	Inclusive of automatic foetal biometry tools measuring key foetal parameters like head circumference (HC), abdominal circumference (AC), femur length (FL) and biparietal diameter (BPD).					
35.7	Inclusive of automated carotid intima-media thickness (CIMT) measurement tools.					
35.8	Support panoramic imaging up to 60cm and curved imaging up to 360°					
35.9	Inclusive of Shear wave elastography (SWE) and advanced strain imaging features for both quantitative and qualitative elastography and to measure tissue deformation in response to compression, ideal for small parts like breast and thyroid.					
35.10	Inclusive of quantitative assessment of liver fat content using ultrasound or ultrasound- derived fat fraction (UDFF).					
36	NETWORKING, CONNECTIVITY AND PACS INTEGRATION					
36.1	Must fully support and comply to DICOM 3.0 standards, including: 1. DICOM Image Storage, 2. DICOM Modality Worklist (MWL), 3. DICOM Query/Retrieve (Q/R) and 4. DICOM Structured Reporting (SR)					
36.2	Must support Modality Performed Procedure Step (MPPS)					
36.4	Allow bidirectional data exchange for seamless workflow.					
36.5	Image must be able to store as DICOM and PC format					

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

	ITEM 1: HIGH-END ULTRASOUND SYSTEM					
	SECTION 1 – USER REQUIREMENTS					
1	1 SYSTEM ARHITECTURE					
REF.	DESCRIPTION	Tick (√)		STATE OR SPECIFY OR REMARKS OR BROCHURE PAGE		
NO.	DESCRIPTION		NO			
36.6	System must have wireless connectivity and ethernet RJ45					
36.7	System must be HL7 ready and activated					
	The systems proposed shall be integrated and connected to Radiology PACS (RPACS) therefore ALL licenses for integration with PACS should be included.					
36.8	Tenderer to include the cost of PACS integration quoted from PACS vendor.					
	Please refer to PACS vendor for information on PACS integration requirement. Tenderer may contact the quotation of PACS integration for this project directly to the MOH's PACS vendor or through the Healthcare Technology Department, Ministry of Health					

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

2	TRANSDUCERS			
REF.	DESCRIPTION	Tick (√)		STATE OR SPECIFY OR REMARKS
NO.	DESCRIPTION	YES	NO	BROCHURE PAGE
	The following transducer shall be included for each of the three high end ultrasound system:			
1	Three (3) units of high frequency linear transducer for vascular, small parts, breast and musculoskeletal. (Tenderer to state the frequency.)			Model: Frequency:
2	Three (3) units of broadband curved array transducer for general purpose abdominal and obstetric. (Tenderer to state the frequency.)			Model: Frequency:
3	Three (3) units of broadband phased array transducer (micro-convex or sector) for cardiac echo and paediatric transcranial ultrasound. (Tenderer to state the frequency.)			Model: Frequency:
4	Three (3) units of broadband curved array transducer for liver shear wave elastography (applicable only if a separate probe from the above is required for this purpose)			Model: Frequency:
5	Three (3) units of linear array hockey stick transducer for peripheral vascular, small parts and superficial structures. (Tenderer to state the frequency.)			Model: Frequency:
6	One (1) unit of curved intra-cavitary transducer for transvaginal obstetrics and gynaecology scan for RIPASH Radiology. Tenderer to state the frequency.			Model: Frequency:
7	All transducers supplied must have a warranty period of at least one (1) year or better and comes with the necessary probe cover (if applicable)			

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

3	DELIVERY AND INSTALLATION SITES			
REF.	DESCRIPTION	Tick	x (✓)	STATE OR SPECIFY OR REMARKS OR
NO.	Y	Y	Ν	BROCHURE PAGE
	Distribution for delivery, installation, testing and commissioning sites are as follows:			
1	One (1) unit for Main ultrasound room, Radiology Department, Raja Isteri Pengiran Anak Saleha (RIPAS) Hospital, Brunei Muara.			
2	One (1) unit for Radiology Unit, Pengiran Muda Mahkota Pengiran Muda Haji Al-Muhtadee Billah (PMMPMHAMB) Hospital, Tutong			
3	One (1) unit for Radiology Unit, Suri Seri Begawan (SSB) Hospital, Kuala Belait			

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

4	WARRANTY AND 3 YEARS POST WARRANTY PREVENTIVE MAINTENANCE (PM)			
REF.	DESCRIPTION	Tick (√)		STATE OR SPECIFY OR REMARKS
NO.	DESCRIPTION	Y	Ν	OR BROCHURE PAGE
1	Tenderer to include warranty period of at least two (2) years			
2	Tenderers to ACKNOWLEDGE the Warranty Undertaking Form in Section 4 stating the terms of warranty provided for the equipment in the tender for the period of two years. This includes but not limited to:			
2.1	Scope of Warranty			
2.2	Two time Planned Preventive Maintenance during warranty [Manufacturer's Standard]			
3	Tenderer to provide additional 3 years Preventive Maintenance service post warranty [At			3 YRS PM POST WARRANTY COST PER UNIT PER YEAR: BND\$
5	least One Time PM per year or as per Manufacturer's Standard]	NUMBER OF PM PER YEAR:	NUMBER OF PM PER YEAR:	
				TOTAL COST FOR 3 YEARS: BND\$
4	Tenderer to provide and attach the scope of work for 3 years PM service post warranty			

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

5	END USER TRAINNING					
REF.	DESCRIPTION	Tick (√)		Tick (√)		STATE OR SPECIFY OR REMARKS OR BROCHURE PAGE
NO.	DESCRIPTION	Y	N			
1	Inclusive of clinical application training for all radiologists for at least seven (7) working days on-site for all the operation and applications offered with the system by Manufacturer's Application Specialist which includes but not limited to:					
1.1	Basic user operation, including image transfers.					
1.2	Application training for the use of the various applications provided with the system.					
1.3	Basic maintenance, including troubleshooting					
2	Training certificate must be provided by manufacturer or tenderer after completion of training sessions.					
3	On-site follow up application training by application specialist after three (3) months of clinical use to ensure the system is fully optimised.					
4	Two (2) sets of User/Operation Manual in English					
5	Two (2) sets of Training Manual in English					
6	Introductory Technical Training to Biomedical Engineers and Technicians at BME RIPASH Office by competent Tenderer's Engineer/Technicians that includes but not limited to:					
6.1	Troubleshooting and basic corrective maintenance					
6.2	Handling and basic inspection maintenance					

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

6	INTERNATIONAL STANDARDS			
REF.	DESCRIPTION	Tick (✔)		STATE OR SPECIFY OR REMARKS OR
NO.	DESCRIPTION	Y	Ν	BROCHURE PAGE
	Equipment model offered must at least be in compliance and approved with four (4) of the following standards or better: (Please provide the documentation/certification to support this compliance)			
1	European Union (CE MARK)			
2	Food and Drug Administration (FDA) USA			
3	International Electrotechnical Commission (IEC) 60601			
4	International Electrotechnical Commission (IEC) 62304			
5	International Organization for Standardization (ISO) 14971			
6	International Organization for Standardization (ISO) 13485			
7	International Organization for Standardization (ISO) 10993			

7	DEMO UNIT			
REF. NO.	DESCRIPTION	Tick	‹ (∕)	STATE OR SPECIFY OR REMARKS OR
	DESCRIPTION	Y	Ν	BROCHURE PAGE
1	Tenderer to arrange a demo unit of the proposed system to be use for at least one to two (2) weeks – in Radiology Department, RIPASH. Please provide details of arranged demo unit			

TENDER FORM

To:

TENDER REFERENCE NO: KK/139/2025/HTD

SECTION 2 – PRICE PROPOSAL		
UNIT PRICE:	TOTAL PRICE:	
BND\$	BND\$	

TENDER FORM

To:

TENDER REFERENCE NO: KK/139/2025/HTD

SECTION 3 - PROCUREMENT AND TECHNICAL SPECIFICATION										
BRAND:						MODEL:				
COUNTRY OF ORIGIN:					UNIT PRICE (B\$):					
WARRANTY PERIOD:						TOTAL PRICE (B\$):				
YEAR INTRODUCED MARKET:	Т	0				LAST COUNTRY SOLD TO:				
PRICE VALIDITY: [AT LEAST ONE (1) Y PRICE VALIDTY]	ΈA	R				DELIVERY TIME:				
AUTHORIZED						APPOINTED BRUN	IEI DIST	RIBUTOR		
AUTHORIZED						PROCURE FROM OVERSEA	COMP	ANY NAME:		
ATTACHED)	IIE	ĸ				AUTHORIZED DISTRIBUTOR	COMP	ANY ORIGIN:		
DETAILED BROCH INCLUDED	IUR	E	YES		NO	⊠ or sp	ecify w	here appropriate		
USER AND SERVICE MANUALS:			YES		NO	Tenderers to acknowledge that they must provide at least TWO sets of USER AND SERVICE manuals when applying commissioning form. One Set for End User, One Set for BME, (Please provide hardcopy or softcopy)				
		22	0V-240	V	BAT	TERY[]YES[]N	0			
MAINS POWER SUPPLY:		5	50-60HZ			of Battery:	Rating:			
		0	THERS	:		RECHARGEABLE		NON-RECHARGEABLE		
POWER ADAPTER/CH OUTPUT RATING:	IAR	GER				EQUIPMENT AMBIENT OPERATING TEMPERATURE RANGE:				
NUMBER OF TEC SUPPORT	CHN	ICAL	LOCA	۱L		 Trained / Certified Not vet trained on the product 				
(ENGINEER/TECHNICI/ Please provide train certification for locals trained/certified DIMENSIONS AND	or Io is	OVEF (SPE) LOCA	RSE CIF` ATIC	A Y DN)	NEAREST LOCATI	ON:	Kilogram (Kg)			
WEIGHT OF MAIN UNIT:						cm inch		Gram(g) Pound (lbs)		
EQUIPMENT WHOLE The supplier shall ensure that spare parts for the equipment are available minimum of 8 years after installation, with the support period extending beyone expected lifecycle of the equipment. No of years: (Please specify)					quipment are available for a period extending beyond the					

TENDER FORM

To:

TENDER REFERENCE NO: KK/139/2025/HTD

INVITATION TO TENDER

PROVISION OF ULTRASOUND SYSTEMS FOR RADIOLOGY DEPARTMENT, MINISTRY OF HEALTH

SECTION 4 – WARRANTY UNDERTAKING FORM

Tenderer, on behalf of the manufacturer, acknowledged and agrees that when equipment is under the warranty period, must cover the scope of normal warranty below at no additional cost:

NORMAL WARRANTY

- Warrants the supplied medical equipment and its accessories to be in good condition, in working order and free from defects to the extend such equipment do not comply with specifications, under normal use for the warranty period. The scope of warranty covers to its maximum extent permitted by applicable law.
- During warranty, tenderer must rectify issues arise from any mechanical, technical or software faulty as soon as it is reported.
- Exchange warranty; Providing replacement units or OEM parts:
 - A. Warranty against defects Manufacturing defects or Equipment malfunction resulted from mechanical, electrical or software failure during Commissioning or within the first _____ months of use
 - B. Faulty workmanship or unsatisfactory condition during delivery or commissioning
 - C. If a unit or accessory is deemed used item or refurbished item (not a new unit) by the user and BME Unit.
- <u>time Planned Preventive Maintenance (PPM) PER YEAR</u> according to Manufacturer's Preventive Maintenance Guideline and to include one-time replacements of battery at the end of 2 years warranty period or any other relevant parts to prolong equipment lifespan.

EXCLUSION FROM WARRANTY

MOH understand that the following circumstances are not covered in the warranty and Tenderer may quote for repair and subject to MOH approval:

- Unauthorized modifications an alteration or repair by anyone other than the Manufacturer or Authorized agent during warranty period.
- Accidental damage or problems caused by negligence or mishandling, subject to appropriate justification by both parties.
- Vandalism and Natural disasters
- Normal wear and tear

ANY OTHER EXCLUSION

Tenderer may propose to include items or terms which is not listed in the exclusion list above for MOH consideration. Please attach behind this form.

TENDERER ACKNOWLEDGMENT

COMPANY CHOP AND SIGNATURE

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

ITEM 2: BREAST IMAGING ULTRASOUND SYSTEM				
	SECTION 1 – USER			
1	SYSTEM ARHITECTURE			
REF.	DESCRIPTION	Tick	κ (√)	STATE OR SPECIFY OR REMARKS
NO.	DESCRIPTION	YES	NO	OR BROCHURE PAGE
1	Two (2) units of cart-based ultrasound (US) system.			
2	Configured specifically for breast and small parts imaging.			
3	The system is fully digital, compact, lightweight and cart-based design			
4	Comes with independent steer and central locking wheels.			
5	Comes with front and rear handles.			
6	Has at least three (3) active transducer ports and one (1) parking probe port.			
7	Has alphanumeric soft keys physical keyboard with easy access scans controls and facility to sanitise the system's keyboard to avoid cross contamination.			
8	Has trackball for moving cursor			
9	Inclusive of two (2) gel bottle holders.			
10	Inclusive of one (1) unit of integrated gel warmer.			
11	Has integrated cable management.			

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

ITEM 2: BREAST IMAGING ULTRASOUND SYSTEM				
	SECTION 1 – USER			
1	SYSTEM ARHITECTURE			
REF.	DESCRIPTION	Tick	κ (√)	STATE OR SPECIFY OR REMARKS
NO.	DESCRIPTION	YES	NO	OR BROCHURE PAGE
12	System storage: integrated SSD or HDD or both.			
13	System storage: at least 1.5TB in total			
14	Operating system: Window 10 or latest			
15	Operating system: install on SSD			
16	Has cybersecurity program to protect system			
17	Power: able to operate on AC and on battery for at least 60 minutes or better			
18	Battery: Self charging and has low-battery indicator			
19	External data interface such as integrated DVD-RW drive, USB or its equivalent to enable external images transfer manually.			
20	Capable to compare images from previous study.			
21	Multiple preloaded application pre-sets available.			
22	Control Monitor/Touch Panel Type: Touchscreen colour display.			

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

ITEM 2: BREAST IMAGING ULTRASOUND SYSTEM							
	SECTION 1 – USER						
1	SYSTEM ARHITECTURE						
REF.	DESCRIPTION	Tick	κ (√)	STATE OR SPECIFY OR REMARKS			
NO.	DESCRIPTION	YES	NO	OR BROCHURE PAGE			
23	Control Monitor/Touch Panel Size: at least 9 inches or better						
24	Control Monitor/Touch Panel has back-lighting						
25	Display Viewing Monitor Type: LCD technology or better						
26	Display Viewing Monitor Size: at least 22" or better						
27	Display Viewing Monitor Resolution: Full HD (1920 x 1080) or better						
28	Display Viewing Monitor has adjustable arm: can adjust tilt, swivel and height.						
29	Display Viewing Monitor has adjustable contrast and brightness						
30	Display Viewing Monitor can be fold down and locked for transportation.						
31	Labelling and markings shall be clear legible and durable. (Durable to withstand routine cleaning)						
32	Can be fold down and locked for transportation.						
33	OPERATING MODES The proposed system must include but not limited to the following operating modes:						

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

ITEM 2: BREAST IMAGING ULTRASOUND SYSTEM				
	SECTION 1 – USER			
1	SYSTEM ARHITECTURE			
REF.	DESCRIPTION	Tick	κ (√)	STATE OR SPECIFY OR REMARKS
NO.	DESCRIPTION	YES	NO	OR BROCHURE PAGE
33.1	Brightness-mode (B) for fundamental imaging mode for anatomical visualization.			
33.2	Motion-mode (M) for motion analysis.			
33.3	Trapezoidal mode for extended field of view for linear probes.			
33.4	Pulsed wave (PW) mode for assessment of blood flow velocities			
33.5	Continuous wave (CW) doppler mode for analysing high-velocity blood flow			
33.6	Colour flow (CF) mode for mapping blood flow direction and velocity.			
33.7	Power doppler imaging (PDI) mode for detecting low-velocity blood flow.			
33.8	Contrast and tissue harmonic imaging for enhancing imaging for vascular studies and for clearer visualization of structures.			
33.9	Spatial compounding for improving image quality by combining multiple beam angles.			
33.10	Extended and convex field of view for enhancing visualization for larger areas during scanning			
33.11	Speckle reduction imaging for reducing noises for clearer images			

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

ITEM 2: BREAST IMAGING ULTRASOUND SYSTEM							
	SECTION 1 – USER						
1	SYSTEM ARHITECTURE						
REF.	DESCRIPTION	Tick	κ (√)	STATE OR SPECIFY OR REMARKS			
NO.	DESCRIPTION	YES	NO	OR BROCHURE PAGE			
33.12	Automatic image optimisation for B-mode and doppler.						
33.13	Adjustment of focal zone, gain, TGC, dynamic range, edge enhancement, gamma correction.						
33.14	Frequency tuning for probes to adjust the frequency range on all transducers.						
33.15	Operator controlled multiple and adjustable focal zones.						
33.16	Real-time pan zoom capability						
33.17	Tools for labelling and anatomical markers for image annotation						
33.18	Customizable user defined pre-sets for variety of clinical applications.						
33.19	2D cine memory and M-mode scroll memory.						
33.20	Cine loop facility up to one minute at 25 frames per second (fps).						
33.21	Fast random-access image review.						
33.22	Ability to apply wide variety of image processing after the examination.						

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

ITEM 2: BREAST IMAGING ULTRASOUND SYSTEM					
	SECTION 1 – USER				
1	SYSTEM ARHITECTURE				
REF.	DESCRIPTION		κ (√)	STATE OR SPECIFY OR REMARKS	
NO.			NO	OR BROCHURE PAGE	
33.23	Full size and split-screen display.				
33.24	Support the following simultaneous display capability: B/PW				
33.25	Support the following simultaneous display capability: B/CF				
33.26	Support the following simultaneous display capability: B/spatial compounding				
33.27	Support the following simultaneous display capability: B/PDI				
33.28	Support the following simultaneous display capability: B/M				
33.29	Multi-image split-screen display capability with combination such as: Live and frozen image comparisons				
33.30	Multi-image split-screen display capability with combination such as: B or spatial compounding and B or spatial compounding				
33.31	Multi-image split-screen display capability with combination such as: B or spatial compounding and CF				
33.32	Multi-image split-screen display capability with combination such as: B or spatial compounding and PDI				
33.33	Multi-image split-screen display capability with combination such as: PW/M				

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

ITEM 2: BREAST IMAGING ULTRASOUND SYSTEM						
	SECTION 1 – USER					
1	SYSTEM ARHITECTURE					
REF.	DESCRIPTION	Tick	κ (√)	STATE OR SPECIFY OR REMARKS		
NO.	DESCRIPTION	YES	NO	OR BROCHURE PAGE		
34	MEASUREMENT AND CALCULATION PACKAGES The proposed system shall include the following:					
34.1	Basic measurements such as: distance (four points), angle (two points), calculation of diameter, area, circumference, calculation of volume by height/width/length, planimetry, automatic planimetry and empirical method.					
34.2	Automated lesion analysis tools: auto lesion outline, volume calculation, BI-RADS Assistant.					
34.3	Real-time automatic doppler and vascular calculations and waveform indices.					
34.4	Elastography Measurements:					
34.4.1	Shear wave elastography (SWE) measurement for breast stiffness in both m/s and kPa with automated interquartile range (IQR) and median value.					
34.4.2	Strain elastography (SE) to evaluate stiffness (elasticity) of breast tissue measure tissue deformation in response to compression and provide a color-coded or greyscale map to represent the stiffness.					
34.5	Biopsy and interventional measurement tools such as needle paths, visibility, distance- to-target measurement, localisation and marking.					
34.6	Inclusive of breast productivity package or breast-specific lesion measurements such as aspect ratio and tumour-to-nipple distance along with tools for evaluating breast productivity.					

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

ITEM 2: BREAST IMAGING ULTRASOUND SYSTEM							
	SECTION 1 – USER						
1	SYSTEM ARHITECTURE						
REF.	DESCRIPTION	Tick	κ (√)	STATE OR SPECIFY OR REMARKS			
NO.	DESCRIPTION	YES	NO	OR BROCHURE PAGE			
34.7	Inclusive of automated carotid intima-media thickness (CIMT) measurement tools.						
34.8	Support panoramic imaging up to 60cm and curved imaging up to 360°						
35	NETWORKING, CONNECTIVITY AND PACS INTEGRATION						
35.1	Must fully support and comply to DICOM 3.0 standards, including: 1. DICOM Image Storage, 2. DICOM Modality Worklist (MWL), 3. DICOM Query/Retrieve (Q/R) and 4. DICOM Structured Reporting (SR)						
35.2	Must support Modality Performed Procedure Step (MPPS)						
35.4	Allow bidirectional data exchange for seamless workflow.						
35.5	Image must be able to store as DICOM and PC format						
35.6	System must have wireless connectivity and ethernet RJ45						
35.7	System must be HL7 ready and activated						
35.8	The systems proposed shall be integrated and connected to Radiology PACS (RPACS) therefore ALL licenses for integration with PACS should be included.						

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

	ITEM 2: BREAST IMAGING ULTRASOUND SYSTEM						
	SECTION 1 – USER						
1	SYSTEM ARHITECTURE						
REF.	DECODIDITION		‹ (✓)	STATE OR SPECIFY OR REMARKS			
NO.	DESCRIPTION	YES	NO	OR BROCHURE PAGE			
	Tenderer to include the cost of PACS integration quoted from PACS vendor.						
	Tenderer may contact the quotation of PACS integration for this project directly to the						
	MOH's PACS vendor or through the Healthcare Technology Department, Ministry of Health						

2	TRANSDUCERS			
REF.	DESCRIPTION	Tick	(✓)	STATE OR SPECIFY OR REMARKS
NO.		YES	NO	OR BROCHURE PAGE
	The following transducers shall be included for each of the two breast imaging ultrasound systems:			
1	Two (2) units of high frequency linear transducer for vascular, small parts, breast and musculoskeletal. (Tenderer to state the frequency.)			Model: Frequency:
2	Two (2) units of broadband curved array transducer for general purpose ultrasound imaging. (Tenderer to state the frequency.)			Model: Frequency:
3	Two (2) units of linear transducer for breast applications. (Tenderer to state the frequency.)			Model:

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

2	TRANSDUCERS			
REF.	DESCRIPTION		(✓)	STATE OR SPECIFY OR REMARKS
NO.	DESCRIPTION			OR BROCHURE PAGE
				Frequency:
4	All transducers supplied must have a warranty period of at least one (1) year or better and comes with the necessary probe cover (if applicable)			

3	DELIVERY AND INSTALLATION SITES			
REF.	DESCRIPTION		(√)	STATE OR SPECIFY OR REMARKS
NO.	DESCRIPTION	YES	NO	OR BROCHURE PAGE
	Distribution for delivery, installation, testing and commissioning sites are as follows:			
1	Two (2) units to Breast Imaging Centre, Raja Isteri Pengiran Anak Saleha (RIPAS) Hospital, Brunei Muara.			

4	WARRANTY AND 3 YEARS POST WARRANTY PREVENTIVE MAINTENANCE (PM)			
REF.	DESCRIPTION	Tick	(√)	STATE OR SPECIFY OR REMARKS OR
NO.	DESCRIPTION	YES	NO	BROCHURE PAGE
1	Tenderer to include warranty period of at least two (2) years			

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

4	WARRANTY AND 3 YEARS POST WARRANTY PREVENTIVE MAINTENANCE (PM)			
REF.		Tick	(✓)	STATE OR SPECIFY OR REMARKS OR
NO.	DESCRIPTION			BROCHURE PAGE
2	Tenderers to ACKNOWLEDGE the Warranty Undertaking Form in Section 4 stating the terms of warranty provided for the equipment in the tender for the period of two years. This includes but not limited to:			
2.1	Scope of Warranty			
2.2	Two time Planned Preventive Maintenance during warranty [Manufacturer's Standard]			
	Tenderer to provide additional 3 years Preventive Maintenance service post warranty [At			3 YRS PM POST WARRANTY COST PER UNIT PER YEAR: BND\$
3	least One Time PM per year or as per Manufacturer's Standard]			NUMBER OF PM PER YEAR:
				TOTAL COST FOR 3 YEARS: BND\$
4	Tenderer to provide and attach the scope of work for 3 years PM service post warranty			

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

5	END USER TRAINNING			
REF.		Tick	(*)	STATE OR SPECIFY OR REMARKS
NO.	DESCRIPTION	YES	NO	OR BROCHURE PAGE
1	Inclusive of clinical application training for all radiologists for at least seven (7) working days on-site for all the operation and applications offered with the system by Manufacturer's Application Specialist which includes but not limited to:			
1.1	Basic user operation, including image transfers.			
1.2	Application training for the use of the various applications provided with the system.			
1.3	Basic maintenance, including troubleshooting			
2	Training certificate must be provided by manufacturer or tenderer after completion of training sessions.			
3	On-site follow up application training by application specialist after three (3) months of clinical use to ensure the system is fully optimised.			
4	Two (2) sets of User/Operation Manual in English			
5	Two (2) sets of Training Manual in English			
6	Introductory Technical Training to Biomedical Engineers and Technicians at BME RIPASH Office by competent Tenderer's Engineer/Technicians that includes but not limited to:			
6.1	Troubleshooting and basic corrective maintenance			
6.2	Handling and basic inspection maintenance			

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

6	INTERNATIONAL STANDARDS			
REF.		Tick	(√)	STATE OR SPECIFY OR REMARKS
NO.	DESCRIPTION		NO	OR BROCHURE PAGE
	Equipment model offered must at least be in compliance and approved with four (4) of the following standards or better: (Please provide the documentation/certification to support this compliance)			
1	European Union (CE MARK)			
2	Food and Drug Administration (FDA) USA			
3	International Electrotechnical Commission (IEC) 60601			
4	International Electrotechnical Commission (IEC) 62304			
5	International Organization for Standardization (ISO) 14971			
6	International Organization for Standardization (ISO) 13485			
7	International Organization for Standardization (ISO) 10993			

7	DEMO UNIT				
REF. NO.	DESCRIPTION			STATE OR SPECIFY OR REMARKS	
	DESCRIPTION	YES	NO	OR BROCHURE PAGE	
1	Tenderer to arrange a demo unit of the proposed system to be use for at least one to two (2) weeks – in Radiology Department, RIPASH.				

TENDER FORM

To:

TENDER REFERENCE NO: KK/139/2025/HTD

SECTION 2 – PRICE PROPOSAL				
UNIT PRICE:	TOTAL PRICE:			
BND\$	BND\$			

TENDER FORM

To:

TENDER REFERENCE NO: KK/139/2025/HTD

SECTION 3 - PROCUREMENT AND TECHNICAL SPECIFICATION											
BRAND:					MODEL	:					
COUNTRY OF ORIGIN:							RICE				
WARRANTY PERIOD:	:					TOTAL	PRICE				
YEAR INTRODUCED MARKET:	Т	0				LAST C SOLD T	OUNTRY O:				
PRICE VALIDITY: [AT LEAST ONE (1) Y PRICE VALIDTY]	ΈA	R				DELIVE TIME:	RY				
						APPOIN		NEI DIS	STRIBUTOR	1	
DISTRIBUTOR:						PROCL	IRE	COMF	PANY NAME	:	
(AUTHORIZED DISTRIBUTOR LET ATTACHED)	ΤE	R				OVERS AUTHO DISTRI	EA RIZED BUTOR	COMPANY ORIGIN:			
DETAILED BROCH INCLUDED	UR	E	YES		NO		⊠ or spe	cify wl	nere approp	oriate	
USER AND SERVICE MANUALS:			YES		NO	Tendere least T when a User, O softcopy	Tenderers to acknowledge that they must provide at least TWO sets of USER AND SERVICE manuals when applying commissioning form. One Set for End User, One Set for BME. (Please provide hardcopy or softcopy)				
		22	0V-240	V	BAT	TERY[]					
MAINS POWER		5	50-60HZ Type			e of Batte	of Battery: Rating:				
		0	THERS	S:		RECHA	RGEABLE		NON-REC	HARGEABLE	
POWER ADAPTER/CHARGER OUTPUT RATING:				·	E OPE	QUIPMENT RATING TE RAN	T AMBI EMPER GE:	ENT ATURE			
NUMBER OF TECI SUPPORT	NICAL LOCAL					□ Tr □ No	ained / Certified				
(ENGINEER/TECHNIC Please provide train certification for locals trained/certified	N) g or no is	OVEF (SPE LOCA	RSE CIF ATIO	EA Y DN)	NEARE	ST LOCAT	ION:				
DIMENSIONS AND					[mm			Kilogram (ł	Kg)	
WEIGHT OF MAIN					[cm			Gram(g))	
UNIT: Image: Display state Image: Display state Pound (lbs) EQUIPMENT The supplier shall ensure that spare parts for the equipment are available for a minimum of 8 years after installation, with the support period extending beyond the expected lifecycle of the equipment. No of years: (Please specify)											

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

INVITATION TO TENDER PROVISION OF ULTRASOUND SYSTEMS FOR RADIOLOGY DEPARTMENT, MINISTRY OF HEALTH

SECTION 4 – WARRANTY UNDERTAKING FORM

Tenderer, on behalf of the manufacturer, acknowledged and agrees that when equipment is under the warranty period, must cover the scope of normal warranty below at no additional cost:

NORMAL WARRANTY

- Warrants the supplied medical equipment and its accessories to be in good condition, in working
 order and free from defects to the extend such equipment do not comply with specifications, under
 normal use for the warranty period. The scope of warranty covers to its maximum extent permitted
 by applicable law.
- During warranty, tenderer must rectify issues arise from any mechanical, technical or software faulty as soon as it is reported.
- Exchange warranty; Providing replacement units or OEM parts:
 - D. Warranty against defects Manufacturing defects or Equipment malfunction resulted from mechanical, electrical or software failure during Commissioning or within the first ______ months of use
 - E. Faulty workmanship or unsatisfactory condition during delivery or commissioning
 - F. If a unit or accessory is deemed used item or refurbished item (not a new unit) by the user and BME Unit.
- <u>time Planned Preventive Maintenance (PPM) PER YEAR</u> according to Manufacturer's Preventive Maintenance Guideline and to include one-time replacements of battery at the end of 2 years warranty period or any other relevant parts to prolong equipment lifespan.

EXCLUSION FROM WARRANTY

MOH understand that the following circumstances are not covered in the warranty and Tenderer may quote for repair and subject to MOH approval:

- Unauthorized modifications an alteration or repair by anyone other than the Manufacturer or Authorized agent during warranty period.
- Accidental damage or problems caused by negligence or mishandling, subject to appropriate justification by both parties.
- Vandalism and Natural disasters
- Normal wear and tear

ANY OTHER EXCLUSION

Tenderer may propose to include items or terms which is not listed in the exclusion list above for MOH consideration. Please attach behind this form.

TENDERER ACKNOWLEDGMENT

COMPANY CHOP AND SIGNATURE

To:

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

	ITEM 3: MID-END ULTRASOUND SYSTEM							
	SECTION 1 – USER REQUIREMENTS							
1	SYSTEM ARHITECTURE							
REF.	DESCRIPTION	Tick	(√)	STATE OR SPECIFY OR				
NO.	DESCRIPTION			PAGE				
1	One (1) unit of cart-based ultrasound (US) system.							
2	Configured specifically for angiography imaging							
3	The system is fully digital, compact, lightweight and easy to move.							
4	Comes with independent steer and central locking wheels.							
5	Comes with front and rear handles.							
6	Has at least two (2) active transducer ports and one (1) parking probe port.							
7	Has alphanumeric soft keys physical keyboard with easy access scans controls and facility to sanitise the system's keyboard to avoid cross contamination.							
8	Has trackball for moving cursor							
9	Inclusive of two (2) gel bottle holders.							
10	Inclusive of one (1) unit of integrated gel warmer.							
11	Has integrated cable management.							

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

ITEM 3: MID-END ULTRASOUND SYSTEM								
	SECTION 1 – USER REQUIREMENTS							
1	SYSTEM ARHITECTURE							
REF.	DESCRIPTION	Tick	(√)	STATE OR SPECIFY OR				
NO.	DESCRIPTION	YES	NO	PAGE				
12	System storage: integrated SSD or HDD or both.							
13	System storage: at least 1.5TB in total							
14	Operating system: Window 10 or latest							
15	Operating system: install on SSD							
16	Has cybersecurity program to protect system							
17	Power: able to operate on AC and on battery for at least 60 minutes or better							
18	Battery: Self charging and has low-battery indicator							
19	External data interface such as integrated DVD-RW drive, USB or its equivalent to enable external images transfer manually.							
20	Capable to compare images from previous study.							
21	Multiple preloaded application pre-sets available.							
22	Control Monitor/Touch Panel Type: Touchscreen colour display.							

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

ITEM 3: MID-END ULTRASOUND SYSTEM							
	SECTION 1 – USER REQUIREMENTS						
1	SYSTEM ARHITECTURE						
REF.		Tick	(√)	STATE OR SPECIFY OR			
NO.	DESCRIPTION		NO	PAGE			
23	Control Monitor/Touch Panel Size: at least 9 inches or better						
24	Control Monitor/Touch Panel has back-lighting						
25	Display Viewing Monitor Type: LCD technology or better						
26	Display Viewing Monitor Size: at least 21" or better						
27	Display Viewing Monitor Resolution: Full HD (1920 x 1080) or better						
28	Display Viewing Monitor has adjustable arm: can adjust tilt, swivel and height.						
29	Display Viewing Monitor has adjustable contrast and brightness						
30	Display Viewing Monitor can be fold down and locked for transportation.						
31	Labelling and markings shall be clear legible and durable. (Durable to withstand routine cleaning)						
32	Can be fold down and locked for transportation.						
33	OPERATING MODES The proposed system must include but not limited to the following operating modes:						

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

ITEM 3: MID-END ULTRASOUND SYSTEM						
SECTION 1 – USER REQUIREMENTS						
1	SYSTEM ARHITECTURE					
REF. NO.	DESCRIPTION	Tick (✓)		STATE OR SPECIFY OR		
		YES	NO	PAGE		
33.1	Brightness-mode (B) for fundamental imaging mode for anatomical visualization.					
33.2	Motion-mode (M) for motion analysis.					
33.3	Trapezoidal mode for extended field of view for linear probes.					
33.4	Pulsed wave (PW) mode for assessment of blood flow velocities					
33.5	Continuous wave (CW) doppler mode for analysing high-velocity blood flow					
33.6	Colour flow (CF) mode for mapping blood flow direction and velocity.					
33.7	Power doppler imaging (PDI) mode for detecting low-velocity blood flow.					
33.8	Contrast and tissue harmonic imaging for enhancing imaging for vascular studies and for clearer visualization of structures.					
33.9	Spatial compounding for improving image quality by combining multiple beam angles.					
33.10	Extended and convex field of view for enhancing visualization for larger areas during scanning					
33.11	Speckle reduction imaging for reducing noises for clearer images					

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

ITEM 3: MID-END ULTRASOUND SYSTEM							
SECTION 1 – USER REQUIREMENTS							
1	SYSTEM ARHITECTURE						
REF. NO.	DESCRIPTION	Tick (√)		STATE OR SPECIFY OR			
		YES	NO	PAGE			
33.12	Automatic image optimisation for B-mode and doppler.						
33.14	Adjustment of focal zone, gain, TGC, dynamic range, edge enhancement, gamma correction.						
33.15	Frequency tuning for probes to adjust the frequency range on all transducers.						
33.16	Operator controlled multiple and adjustable focal zones.						
33.17	Real-time pan zoom capability						
33.18	Tools for labelling and anatomical markers for image annotation						
33.19	Customizable user defined pre-sets for variety of clinical applications.						
33.20	2D cine memory and M-mode scroll memory.						
33.21	Cine loop facility up to one minute at 25 frames per second (fps).						
33.22	Fast random-access image review.						
33.23	Ability to apply wide variety of image processing after the examination.						

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

ITEM 3: MID-END ULTRASOUND SYSTEM						
SECTION 1 – USER REQUIREMENTS						
1	SYSTEM ARHITECTURE					
REF. NO.	DESCRIPTION	Tick (✔)		STATE OR SPECIFY OR		
		YES	NO	PAGE		
33.24	Full size and split-screen display.					
33.25	Support the following simultaneous display capability: B/PW					
33.26	Support the following simultaneous display capability: B/CF					
33.27	Support the following simultaneous display capability: B/spatial compounding					
33.28	Support the following simultaneous display capability: B/PDI					
33.29	Support the following simultaneous display capability: B/M					
33.30	Multi-image split-screen display capability with combination such as: Live and frozen image comparisons					
33.31	Multi-image split-screen display capability with combination such as: B or spatial compounding and B or spatial compounding					
33.32	Multi-image split-screen display capability with combination such as: B or spatial compounding and CF					
33.33	Multi-image split-screen display capability with combination such as: B or spatial compounding and PDI					
33.34	Multi-image split-screen display capability with combination such as: PW/M					
TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

	ITEM 3: MID-END ULTRASOUND SYSTEM									
	SECTION 1 – USER REQUIREMENTS									
1	1 SYSTEM ARHITECTURE									
REF.	DECODIDITION	Tick	(√)	STATE OR SPECIFY OR						
NO.	DESCRIPTION	YES	NO	PAGE						
34	MEASUREMENT AND CALCULATION PACKAGES The proposed system shall include the following:									
34.1	Basic measurements such as: distance (four points), angle (two points), calculation of diameter, area, circumference, calculation of volume by height/width/length, planimetry, automatic planimetry and empirical method.									
34.2	Automated measurement packages including reports, urology, renal, cardiac (M-mode and B-mode), obstetrics and gynaecological and vascular.									
34.3	Real-time automatic doppler calculations and waveform indices.									
34.4	Support panoramic imaging up to 60cm and curved imaging up to 360 degrees									
35	NETWORKING, CONNECTIVITY AND PACS INTEGRATION									
35.1	Must fully support and comply to DICOM 3.0 standards, including: 1. DICOM Image Storage, 2. DICOM Modality Worklist (MWL), 3. DICOM Query/Retrieve (Q/R) and 4. DICOM Structured Reporting (SR)									
35.2	Must support Modality Performed Procedure Step (MPPS)									
35.3	Allow bidirectional data exchange for seamless workflow.									

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

	ITEM 3: MID-END ULTRASOUND SYSTEM							
	SECTION 1 – USER REQUIREMENTS							
1	SYSTEM ARHITECTURE							
REF.	DESCRIPTION	Tick	(•⁄)	STATE OR SPECIFY OR				
NO. DESCRIPTION	DESCRIPTION	YES	NO	PAGE				
35.4	Image must be able to store as DICOM and PC format							
35.5	System must have wireless connectivity and ethernet RJ45							
35.6	System must be HL7 ready and activated							
	The systems proposed shall be integrated and connected to Radiology PACS (RPACS) therefore ALL licenses for integration with PACS should be included.							
36.7	Tenderer to include the cost of PACS integration quoted from PACS vendor.							
	Please refer to PACS vendor for information on PACS integration requirement. Tenderer may contact the quotation of PACS integration for this project directly to the MOH's PACS vendor or through the Healthcare Technology Department, Ministry of Health							

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

2	TRANSDUCERS			
REF.	DECODIDITION			STATE OR SPECIFY OR
NO.	DESCRIPTION	YES	NO	PAGE
	The following transducers shall be included for the mid-end ultrasound system:			
1	One (1) units of high frequency linear transducer for vascular, small parts, breast and musculoskeletal. (Tenderer to state the frequency.)			Model: Frequency:
2	One (1) units of broadband curved array transducer for general purpose abdominal and obstetric. (Tenderer to state the frequency.)			Model: Frequency:
3	All transducers supplied must have a warranty period of at least one (1) year or better and comes with the necessary probe cover (if applicable)			

3	DELIVERY AND INSTALLATION SITES			
REF. NO.	DESCRIPTION	Tick	x (✓)	STATE OR SPECIFY OR REMARKS OR BROCHURE PAGE
	DESCRIPTION	YES	NO	
	Distribution for delivery, installation, testing and commissioning sites are as follows:			
1	One (1) unit for Angiography Room, Radiology Department, Raja Isteri Pengiran Anak Saleha (RIPAS) Hospital, Brunei Muara.			

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

4	WARRANTY AND 3 YEARS POST WARRANTY PREVENTIVE MAINTENANCE (PM)			
REF.	RECORDETION	Tick (√)		STATE OR SPECIFY OR
NO.	DESCRIPTION		NO	PAGE
1	Tenderer to include warranty period of at least two (2) years			
2	Tenderers to ACKNOWLEDGE the Warranty Undertaking Form in Section 4 stating the terms of warranty provided for the equipment in the tender for the period of two years. This includes but not limited to:			
2.1	Scope of Warranty			
2.2	Two time Planned Preventive Maintenance during warranty [Manufacturer's Standard]			
				3 YRS PM POST WARRANTY COST PER UNIT PER YEAR: BND\$
3	Time PM per year or as per Manufacturer's Standard]			NUMBER OF PM PER YEAR:
				TOTAL COST FOR 3 YEARS: BND\$
4	Tenderer to provide and attach the scope of work for 3 years PM service post warranty			

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

5	END USER TRAINNING				
REF.	RECORDETION		(~)	STATE OR SPECIFY OR	
NO.	DESCRIPTION	YES NO		PAGE	
1	Inclusive of clinical application training for all radiologists for at least seven (7) working days on- site for all the operation and applications offered with the system by Manufacturer's Application Specialist which includes but not limited to:				
1.1	Basic user operation, including image transfers.				
1.2	Application training for the use of the various applications provided with the system.				
1.3	Basic maintenance, including troubleshooting				
2	Training certificate must be provided by manufacturer or tenderer after completion of training sessions.				
3	On-site follow up application training by application specialist after three (3) months of clinical use to ensure the system is fully optimised.				
4	Two (2) sets of User/Operation Manual in English				
5	Two (2) sets of Training Manual in English				
6	Introductory Technical Training to Biomedical Engineers and Technicians at BME RIPASH Office by competent Tenderer's Engineer/Technicians that includes but not limited to:				
6.1	Troubleshooting and basic corrective maintenance				
6.2	Handling and basic inspection maintenance				

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

6	INTERNATIONAL STANDARDS			
REF.	DESCRIPTION		(✓)	STATE OR SPECIFY OR
NO.			NO	PAGE
	Equipment model offered must at least be in compliance and approved with four (4) of the following standards or better: (Please provide the documentation/certification to support this compliance)			
1	European Union (CE MARK)			
2	Food and Drug Administration (FDA) USA			
3	International Electrotechnical Commission (IEC) 60601			
4	International Electrotechnical Commission (IEC) 62304			
5	International Organization for Standardization (ISO) 14971			
6	International Organization for Standardization (ISO) 13485			
7	International Organization for Standardization (ISO) 10993			

7	DEMO UNIT			
REF. NO.	DESCRIPTION	Tick	(√)	STATE OR SPECIFY OR
		YES	NO	PAGE
1	Tenderer to arrange a demo unit of the proposed system to be use for at least one to two (2) weeks – in Radiology Department, RIPASH.			

TENDER FORM

To:

TENDER REFERENCE NO: KK/139/2025/HTD

SECTION 2 – PR	RICE PROPOSAL
UNIT PRICE:	TOTAL PRICE:
BND\$	BND\$

TENDER FORM

To:

TENDER REFERENCE NO: KK/139/2025/HTD

SECTION 3 - PROCUREMENT AND TECHNICAL SPECIFICATION												
BRAND:						MODEL	:					
COUNTRY OF ORIGIN					UNIT P	RICE						
	:					(B\$): TOTAL (B\$) [:]	PRICE					
YEAR INTRODUCED MARKET:	то					LAST C SOLD 1	OUNTRY O:					
PRICE VALIDITY: [AT LEAST ONE (1) Y PRICE VALIDTY]	EAR					DELIVE TIME:	RY					
AUTHORIZED						APPOIN	NTED BRUI	NEI DIS	STRIBUTOR			
DISTRIBUTOR:						PROCL	IRE	COMF	PANY NAME	:		
(AUTHORIZED DISTRIBUTOR LET ATTACHED)	TER					FROM OVERSEA AUTHORIZED DISTRIBUTOR		COMPANY ORIGIN:				
DETAILED BROCH INCLUDED	URE		YES		NO		⊠ or spe	ecify where appropriate				
USER AND SER' MANUALS:	VICE		YES		NO	Tendere least T when a User, O softcopy	ers to ackno NO sets o oplying com ne Set for l /)	nowledge that they must provide at of USER AND SERVICE manuals ommissioning form. One Set for End r BME. (Please provide hardcopy or				
		22	:0V-240)V	BAT	TERY[]YES[]NO						
		5	0-60HZ	Z	Туре	e of Batte	ry:	Rating:				
		0	THER	S:		RECHA	RGEABLE		NON-REC	HARGEABLE		
POWER ADAPTER/CHARGER OUTPUT RATING:						E OPE	QUIPMEN RATING TI RAN	IT AMBIENT EMPERATURE NGE:				
NUMBER OF TEC	HNIC/	۹L	LOCA	۱L				rained / Certified				
(ENGINEER/TECHNIC Please provide train certification for locals trained/certified	OVERSEA or (SPECIFY is LOCATION)				NEARE	ST LOCAT	ION:					
DIMENSIONS AND					[mm			Kilogram (ł	≺ g)		
WEIGHT OF MAIN							Gram(g)	١				
	The	sur	plier sl	hall	ensur	e that so:	are parts fo	r the ec	uipment are	<i>)</i> e available for a		
EQUIPMENT WHOLE LIFE TIME SUPPORT: The supplier shall ensure that spare parts for the equipment are available for a minimum of 8 years after installation, with the support period extending beyond the expected lifecycle of the equipment. No of years: (Please specify)									tending beyond			

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

INVITATION TO TENDER PROVISION OF ULTRASOUND SYSTEMS FOR RADIOLOGY DEPARTMENT, MINISTRY OF HEALTH

SECTION 4 – WARRANTY UNDERTAKING FORM

Tenderer, on behalf of the manufacturer, acknowledged and agrees that when equipment is under the warranty period, must cover the scope of normal warranty below at no additional cost:

NORMAL WARRANTY

- Warrants the supplied medical equipment and its accessories to be in good condition, in working
 order and free from defects to the extend such equipment do not comply with specifications, under
 normal use for the warranty period. The scope of warranty covers to its maximum extent permitted
 by applicable law.
- During warranty, tenderer must rectify issues arise from any mechanical, technical or software faulty as soon as it is reported.
- Exchange warranty; Providing replacement units or OEM parts:
 - G. Warranty against defects Manufacturing defects or Equipment malfunction resulted from mechanical, electrical or software failure during Commissioning or within the first ______ months of use
 - H. Faulty workmanship or unsatisfactory condition during delivery or commissioning
 - I. If a unit or accessory is deemed used item or refurbished item (not a new unit) by the user and BME Unit.
- <u>time Planned Preventive Maintenance (PPM) PER YEAR</u> according to Manufacturer's Preventive Maintenance Guideline and to include one-time replacements of battery at the end of 2 years warranty period or any other relevant parts to prolong equipment lifespan.

EXCLUSION FROM WARRANTY

MOH understand that the following circumstances are not covered in the warranty and Tenderer may quote for repair and subject to MOH approval:

- Unauthorized modifications an alteration or repair by anyone other than the Manufacturer or Authorized agent during warranty period.
- Accidental damage or problems caused by negligence or mishandling, subject to appropriate justification by both parties.
- Vandalism and Natural disasters
- Normal wear and tear

ANY OTHER EXCLUSION

Tenderer may propose to include items or terms which is not listed in the exclusion list above for MOH consideration. Please attach behind this form.

TENDERER ACKNOWLEDGMENT

COMPANY CHOP AND SIGNATURE

To:

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

ITEM 4: O&G ULTRASOUND SYSTEM								
	SECTION 1 – USER REQUIREMENTS							
1	SYSTEM ARHITECTURE							
REF.	DESCRIPTION	Tic	k (√)	STATE OR SPECIFY OR				
NO.	DESCRIPTION	YES	NO	PAGE				
1	Real-time image optimization and transducer technology migration							
2	Imaging frequency range: 1–20 MHz							
3	At least 4 active transducer ports with electronic selection							
4	Integrated gel warmer with temperature adjustment							
5	Transducer storage holder for minimum 7 probes & utility storage baskets with gel holder							
6	Integrated cable management with multiple cable hooks, front/back handles, and lockable wheels							
7	Customizable imaging modes and parameters							
8	IMAGING CAPABILITIES							
8.1	Imaging modes: B-mode, M-mode, Color & Power Doppler, Pulsed Wave & Continuous Wave Doppler							
8.2	Adjustable field of view size and position, 2D beam steering angle $\geq \pm 30^{\circ}$							
8.3	Parenchymal image harmonization for reduced noise and improved sharpness							

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

ITEM 4: O&G ULTRASOUND SYSTEM								
	SECTION 1 – USER REQUIREMENTS							
1	SYSTEM ARHITECTURE							
REF.	DESCRIPTION	Tic	k (√)	STATE OR SPECIFY OR				
NO.	DESCRIPTION	YES	NO	PAGE				
8.4	Speckle noise reduction with ≥ 3 levels via dynamic tissue contrast enhancement							
9	CONTROL PANEL & DISPLAY	•						
9.1	Height-adjustable control panel with ±90° swivel articulation							
9.2	Tiltable touchscreen interface with USB ports							
9.3	Conventional keyboard and laser optical trackball with backlighting							
9.4	In-Plane Switching monitor							
9.5	Display size: ≥ 21 inches							
9.6	Display resolution ≥ 1920×1080							
9.7	Digital calipers: minimum 4 active							
9.8	Zoom and pan on both real-time and frozen images							
10	POWER, SOFTWARE & STORAGE							

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

ITEM 4: O&G ULTRASOUND SYSTEM									
	SECTION 1 – USER REQUIREMENTS								
1	SYSTEM ARHITECTURE								
REF.	DESCRIPTION	Tic	k (√)	STATE OR SPECIFY OR					
NO.	DESCRIPTION	YES	NO	PAGE					
10.1	Mains supply: 100–240V AC, 50/60 Hz								
10.2	Battery integrated								
10.3	Battery operation: ≥ 70 minutes scan time with status display								
10.4	Operating System: Windows 10 or later with antivirus installed								
10.5	SSD ≥ 500 GB with automatic disk management								
10.6	Cine memory: not less than 500 frames								
11	NETWORKING, CONNECTIVITY AND PACS INTEGRATION								
11.1	Must fully support and comply to DICOM 3.0 standards, including: 1. DICOM Image Storage, 2. DICOM Modality Worklist (MWL), 3. DICOM Query/Retrieve (Q/R) and 4. DICOM Structured Reporting (SR)								
11.2	Must support Modality Performed Procedure Step (MPPS)								
11.3	Allow bidirectional data exchange for seamless workflow.								

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

	ITEM 4: O&G ULTRASOUND SYSTEM									
	SECTION 1 – USER REQUIREMENTS									
1	SYSTEM ARHITECTURE									
REF.	DESCRIPTION	Ticł	‹ (✓)	STATE OR SPECIFY OR						
NO.	DESCRIPTION	YES	NO	PAGE						
11.4	Image must be able to store as DICOM and PC format									
11.5	System must have wireless connectivity and ethernet RJ45									
11.6	System must be HL7 ready and activated									
	The systems proposed shall be integrated and connected to Radiology PACS (RPACS) therefore ALL licenses for integration with PACS should be included.									
11.7	Tenderer to include the cost of PACS integration quoted from PACS vendor.									
	Please refer to PACS vendor for information on PACS integration requirement. Tenderer may contact the quotation of PACS integration for this project directly to the MOH's PACS vendor or through the Healthcare Technology Department, Ministry of Health									

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

2	TRANSDUCERS				
REF.	DESCRIPTION			STATE OR SPECIFY OR	
NO.	DESCRIPTION		NO		AGE
	The following transducers shall be included for the O&G ultrasound system:				
1	One (1) unit of multi-frequency convex probe Penetration \ge 350 mm and Field of View \ge 70 ⁰			Model:	FOV:
	(Tenderer to state the Model, Penetration, FOV and Frequency)			Penetration:	Frequency:
	One (1) unit of Endo Cavity Transducer			Model:	FOV:
2	■ penetration ≥ 140 mm and Field of View ≥ 220 ⁰				
	(Tenderer to state the Model, Penetration, FOV and Frequency)			Penetration:	Frequency:
	One (1) unit of 4D Volume Transducer			Model:	FOV:
3	■ penetration ≥ 300 mm and Field of View ≥ 65 ⁰				
	(Tenderer to state the Model, Penetration, FOV and Frequency)			Penetration:	Frequency:
	One (1) unit of 4D Endo Cavity Volume Transducer			Model:	FOV:
4	■ penetration ≥ 160 mm and Field of View ≥ 145 ^o				
	(Tenderer to state the Model, Penetration, FOV and Frequency)			Penetration:	Frequency:
5	All transducers supplied must have a warranty period of at least one (1) year or better and comes				
Э	with the necessary probe cover (if applicable)				

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

3	DELIVERY AND INSTALLATION SITES				
REF.	DESCRIPTION	Tick (√)		STATE OR SPECIFY OR	
NO.	DESCRIPTION	YES	NO	PAGE	
	Distribution for delivery, installation, testing and commissioning sites are as follows:				
1	One (1) unit for Pengkalan Batu Health Centre, MCH, Brunei Muara.				
2	One (1) unit for Sg Kelugos Health Centre, MCH, Tutong.				
3	One (1) unit for Telisai Health Centre, MCH, Tutong.				

4	WARRANTY AND 3 YEARS POST WARRANTY PREVENTIVE MAINTENANCE (PM)									
REF.	DECODIDITION			STATE OR SPECIFY OR						
NO.	DESCRIPTION	YES	NO	PAGE						
1	Tenderer to include warranty period of at least two (2) years									
2	Tenderers to ACKNOWLEDGE the Warranty Undertaking Form in Section 4 stating the terms of warranty provided for the equipment in the tender for the period of two years. This includes but not limited to:									
2.1	Scope of Warranty									
2.2	Two time Planned Preventive Maintenance during warranty [Manufacturer's Standard]									

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

4	WARRANTY AND 3 YEARS POST WARRANTY PREVENTIVE MAINTENANCE (PM)									
REF. NO.	DESCRIPTION	Tick (✔)		STATE OR SPECIFY OR						
	DESCRIPTION	YES	NO	PAGE						
3	Tenderer to provide additional 3 years Preventive Maintenance service post warranty [At least One Time PM per year or as per Manufacturer's Standard]			3 YRS PM POST WARRANTY COST PER UNIT PER YEAR: BND\$ NUMBER OF PM PER YEAR: TOTAL COST FOR 3 YEARS: BND\$						
4	Tenderer to provide and attach the scope of work for 3 years PM service post warranty									

5	END USER TRAINNING				
REF.	DESCRIPTION		(✓)	STATE OR SPECIFY OR	
NO.	DESCRIPTION	YES	NO	PAGE	
1	Inclusive of clinical application training for all radiologists for at least seven (7) working days on- site for all the operation and applications offered with the system by Manufacturer's Application Specialist which includes but not limited to:				
1.1	Basic user operation, including image transfers.				
1.2	Application training for the use of the various applications provided with the system.				
1.3	Basic maintenance, including troubleshooting				

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

5	END USER TRAINNING			
REF.	DESCRIPTION		(~)	STATE OR SPECIFY OR
NO.			NO	PAGE
2	Training certificate must be provided by manufacturer or tenderer after completion of training sessions.			
3	On-site follow up application training by application specialist after three (3) months of clinical use to ensure the system is fully optimised.			
4	Two (2) sets of User/Operation Manual in English			
5	Two (2) sets of Training Manual in English			
6	Introductory Technical Training to Biomedical Engineers and Technicians at BME RIPASH Office by competent Tenderer's Engineer/Technicians that includes but not limited to:			
6.1	Troubleshooting and basic corrective maintenance			
6.2	Handling and basic inspection maintenance			

6	INTERNATIONAL STANDARDS			
REF. NO.	DESCRIPTION	Tick	(√)	STATE OR SPECIFY OR REMARKS OR BROCHURE PAGE
	DESCRIPTION	YES	NO	
	Equipment model offered must at least be in compliance and approved with four (4) of the following standards or better: (Please provide the documentation/certification to support this compliance)			
1	European Union (CE MARK)			

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

6	INTERNATIONAL STANDARDS				
REF.	DESCRIPTION		(*)	STATE OR SPECIFY OR	
NO.	DESCRIPTION	YES	NO	PAGE	
2	Food and Drug Administration (FDA) USA				
3	International Electrotechnical Commission (IEC) 60601				
4	International Electrotechnical Commission (IEC) 62304				
5	International Organization for Standardization (ISO) 14971				
6	International Organization for Standardization (ISO) 13485				
7	International Organization for Standardization (ISO) 10993				

SECTION 2 – PRICE PROPOSAL							
UNIT PRICE:	TOTAL PRICE:						
BND\$	BND\$						

TENDER FORM

To:

TENDER REFERENCE NO: KK/139/2025/HTD

SECTION 3 - PROCUREMENT AND TECHNICAL SPECIFICATION													
BRAND:					MODEL	:							
COUNTRY OF ORIGIN				UNIT P	RICE								
WARRANTY PERIOD:					(B\$): TOTAL	PRICE							
	то					(B\$):							
MARKET:	10					SOLD T	0:						
PRICE VALIDITY: [AT LEAST ONE (1) Y PRICE VALIDTY]	EAR					DELIVE TIME:	RY						
AUTHORIZED						APPOIN	NTED BRUI	NEI DIS	STRIBUTOR				
DISTRIBUTOR:						PROCL	IRE	COMF	PANY NAME	E			
(AUTHORIZED DISTRIBUTOR LET ATTACHED)	TER					FROM OVERSEA AUTHORIZED DISTRIBUTOR		COMPANY ORIGIN:					
DETAILED BROCH INCLUDED	URE		YES		NO		⊠ or spe	ecify where appropriate					
USER AND SER MANUALS:		YES		NO	Tendere least T when a User, O softcopy	ers to ackno NO sets o oplying com ne Set for l /)	nowledge that they must provide at of USER AND SERVICE manuals ommissioning form. One Set for End r BME. (Please provide hardcopy or						
		22	:0V-240	V	BAT	TERY[]							
		50-60HZ			Туре	of Batte	ry:	Rating:					
SOLLET.		0	THER	S:		RECHARGEABLE NON-R			NON-REC	HARGEABLE			
POWER ADAPTER/CHARGER OUTPUT RATING:					E OPE	QUIPMEN RATING TI RAN	ENT ATURE						
NUMBER OF TECH SUPPORT	۹L	LOCA	۱L				rained / Certified ot yet trained on the product						
(ENGINEER/TECHNIC Please provide train certification for locals trained/certified	OVERSEA or (SPECIFY is LOCATION)				NEARE	ST LOCAT	ION:						
DIMENSIONS AND					[mm			Kilogram (ł	≺g)			
WEIGHT OF MAIN				[cm			Gram(g)	`				
UNIT.	The	sur	oplier sl	hall	ensur	Inch e that spa	are parts fo	r the ec	Pound (IDS) e available for a			
EQUIPMENT WHOLE LIFE TIME SUPPORT: (Please specify)								tending beyond					

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

INVITATION TO TENDER PROVISION OF ULTRASOUND SYSTEMS FOR RADIOLOGY DEPARTMENT, MINISTRY OF HEALTH

SECTION 4 – WARRANTY UNDERTAKING FORM

Tenderer, on behalf of the manufacturer, acknowledged and agrees that when equipment is under the warranty period, must cover the scope of normal warranty below at no additional cost:

NORMAL WARRANTY

- Warrants the supplied medical equipment and its accessories to be in good condition, in working
 order and free from defects to the extend such equipment do not comply with specifications, under
 normal use for the warranty period. The scope of warranty covers to its maximum extent permitted
 by applicable law.
- During warranty, tenderer must rectify issues arise from any mechanical, technical or software faulty as soon as it is reported.
- Exchange warranty; Providing replacement units or OEM parts:
 - J. Warranty against defects Manufacturing defects or Equipment malfunction resulted from mechanical, electrical or software failure during Commissioning or within the first ______ months of use
 - K. Faulty workmanship or unsatisfactory condition during delivery or commissioning
 - L. If a unit or accessory is deemed used item or refurbished item (not a new unit) by the user and BME Unit.
- <u>time Planned Preventive Maintenance (PPM) PER YEAR</u> according to Manufacturer's Preventive Maintenance Guideline and to include one-time replacements of battery at the end of 2 years warranty period or any other relevant parts to prolong equipment lifespan.

EXCLUSION FROM WARRANTY

MOH understand that the following circumstances are not covered in the warranty and Tenderer may quote for repair and subject to MOH approval:

- Unauthorized modifications an alteration or repair by anyone other than the Manufacturer or Authorized agent during warranty period.
- Accidental damage or problems caused by negligence or mishandling, subject to appropriate justification by both parties.
- Vandalism and Natural disasters
- Normal wear and tear

ANY OTHER EXCLUSION

Tenderer may propose to include items or terms which is not listed in the exclusion list above for MOH consideration. Please attach behind this form.

TENDERER ACKNOWLEDGMENT

COMPANY CHOP AND SIGNATURE

To:

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

ITEM 5: ULTRASOUND EXAMINATION COUCH AND CHAIR								
SECTION 1 – USER REQUIREMENTS								
1	SYSTEM ARHITECTURE							
REF.	DESCRIPTION	Tick (√)		STATE OR SPECIFY OR				
NO.	DESCRIPTION	YES	NO	PAGE				
1	Six (6) units of ultrasound examination couch and Ultrasound operator chair							
2	Ultrasound Examination Couch: Electric height adjustable type							
3	Ultrasound Examination Couch: Adjustable back and leg rest.							
4	Ultrasound Examination Couch: Complete with disposable paper dispenser mounts to the head end of the couch.							
5	Ultrasound Examination Couch: Suitable dimension for dedicated room (Please refer to each location)							
6	Ultrasound Chair: Height adjustable ultrasound chair (saddle type) for the operators with wheels and adjustable back							

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

2	DELIVERY AND INSTALLATION SITES							
REF. NO.	DESCRIPTION	Tick	(*)	STATE OR SPECIFY OR REMARKS				
	DESCRIPTION	YES	NO	OR BROCHURE PAGE				
	Distribution for delivery, installation, testing and commissioning sites are as follows:							
1	One (1) unit for Main ultrasound room, Radiology Department, Raja Isteri Pengiran Anak Saleha (RIPAS) Hospital, Brunei Muara.							
2	One (1) unit for Radiology Unit, Pengiran Muda Mahkota Pengiran Muda Haji Al- Muhtadee Billah (PMMPMHAMB) Hospital, Tutong							
3	One (1) unit for Radiology Unit, Suri Seri Begawan (SSB) Hospital, Kuala Belait							
4	One (1) unit for Angio Room, Radiology Department, Raja Isteri Pengiran Anak Saleha (RIPAS) Hospital, Brunei Muara.							
5	Two (2) units for Breast Imaging Centre, Raja Isteri Pengiran Anak Saleha (RIPAS) Hospital, Brunei Muara.							

3	WARRANTY			
REF. NO.	DESCRIPTION			STATE OR SPECIFY OR REMARKS OR
	DESCRIPTION	Y	N	BROCHURE PAGE
1	Tenderer to include warranty period of at least two (2) years			
2	Tenderers to ACKNOWLEDGE the Warranty Undertaking Form in Section 4 stating the terms and scope of warranty provided for the equipment in the tender for the period of two years.			

TENDER FORM

To:

TENDER REFERENCE NO: KK/139/2025/HTD

SECTION 2 – PRICE PROPOSAL							
UNIT PRICE:	TOTAL PRICE:						
BND\$	BND\$						

TENDER FORM

To:

TENDER REFERENCE NO: KK/139/2025/HTD

SECTION 3 - PROCUREMENT AND TECHNICAL SPECIFICATION											
BRAND:						MODEL	MODEL:				
COUNTRY OF ORIGIN						UNIT P	RICE				
WARRANTY PERIOD:						(B\$): TOTAL PRICE (B\$):					
YEAR INTRODUCED TO MARKET:						LAST C SOLD T	OUNTRY O:				
PRICE VALIDITY: [AT LEAST ONE (1) YEAR PRICE VALIDTY]						DELIVE TIME:	RY				
						APPOIN	APPOINTED BRUNEI DISTRIBUTOR				
DISTRIBUTOR:						PROCURE		COMPANY NAME:			
(AUTHORIZED DISTRIBUTOR LETTER ATTACHED)						FROM OVERSEA AUTHORIZED DISTRIBUTOR		COMPANY ORIGIN:			
DETAILED BROCH INCLUDED		YES		NO		⊠ or spe	pecify where appropriate				
USER AND SERVICE MANUALS:			YES		NO	Tenderers to acknowledge that they must provide at least TWO sets of USER AND SERVICE manuals when applying commissioning form. One Set for Enc User, One Set for BME. (Please provide hardcopy of softcopy)					
			0V-240)V	BAT	TERY[]YES[]NO					
	5 C		0-60HZ	Z	Туре	Type of Battery:			Rating:		
SUFFEI.			THERS	S:		RECHA	RGEABLE		NON-REC	HARGEABLE	
POWER ADAPTER/CHARGER OUTPUT RATING:						EQUIPMENT AMBIENT OPERATING TEMPERATURE RANGE:					
NUMBER OF TECHNICA SUPPORT (ENGINEER/TECHNICIAN) Please provide training certification for locals who trained/certified			CAL LOCAL				□ Tr □ No	rained / Certified ot yet trained on the product			
			OVEF (SPEC LOCA	RSE CIF	A Y DN)	NEAREST LOCATION:					
DIMENSIONS AND					[🗆 mm 🛛 🔅 Kilogram (Kg)					
WEIGHT OF MAIN						□ cm □ Gram(g))	
EQUIPMENT WHOLE LIFE TIME SUPPORT: The supplier shall ensure that spare parts for the equipment are available for a minimum of 8 years after installation, with the support period extending beyond the expected lifecycle of the equipment. No of years: (Please specify)											

TENDER FORM

TENDER REFERENCE NO: KK/139/2025/HTD

INVITATION TO TENDER

PROVISION OF ULTRASOUND SYSTEMS FOR RADIOLOGY DEPARTMENT, MINISTRY OF

HEALTH

SECTION 4 – WARRANTY UNDERTAKING FORM

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 normal use for the warranty period. The scope of warranty covers to its maximum extent permitted
 by applicable law.
- During warranty, tenderer must rectify issues arise from any mechanical, technical or software faulty as soon as it is reported.
- Exchange warranty; Providing replacement units or OEM parts:
 - M. Warranty against defects Manufacturing defects or Equipment malfunction resulted from mechanical, electrical or software failure during Commissioning or within the first ______ months of use
 - N. Faulty workmanship or unsatisfactory condition during delivery or commissioning
 - O. If a unit or accessory is deemed used item or refurbished item (not a new unit) by the user and BME Unit.
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- Normal wear and tear

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TENDERER ACKNOWLEDGMENT

COMPANY CHOP AND SIGNATURE

To: