

**TENDER REFERENCE NO.: KK/101/2026/HTD**

**MINISTRY OF HEALTH  
NEGARA BRUNEI DARUSSALAM**

**SUPPLY, DELIVERY, INSTALLATION, TESTING,  
COMMISSIONING AND UPGRADE OF EXISTING SURGICAL  
ENDOSCOPY CAMERA HEAD SYSTEM FOR OPERATING  
THEATRE, RIPAS HOSPITAL**

**TENDER FEES : \$10.00**

**RECEIPT NO. :**

**CLOSING DATE : ON Tuesday, 09th June 2026**

**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**

**(CLUSTERING)**

**SECTION 2**

**SPECIFICATIONS AND REQUIREMENTS**

**TENDER REFERENCE NO: KK/101/2026/HTD**

**INVITATION TO TENDER**

**SUPPLY, DELIVERY, INSTALLATION, TESTING, COMISSIONING AND UPGRADE OF EXISTING  
SURGICAL ENDOSCOPY CAMERA HEAD SYSTEM FOR OPERATING THEATRE, RIPAS  
HOSPITAL**

<b>DELIVERY PERIOD</b>	<b>NOT MORE THAN 90 DAYS UPON CONFIRMATION</b>
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<b>SCOPE OF WORK AND SUMMARY OF PRICES</b>
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**This tender is for the Supply, Delivery, Installation, Testing, Commissioning, Warranty and Maintenance of:**

<b>DESCRIPTION</b>	<b>QUANTITY</b>
<b>UPGRADE OF EXISTING SURGICAL ENDOSCOPY CAMERA HEAD SYSTEM</b>	<b>2</b>

SECTION 1 – USER REQUIREMENTS														
REF. NO.	DESCRIPTION													
1	<b>SYSTEM ARCHITECTURE</b>													
1.1	<p>The upgrade shall enhance an <b>existing Karl Storz surgical endoscopy system</b> installed in the operating theatre, to support <b>laparoscopic and endoscopic procedures for General Surgery and Gynaecology</b>, without replacement of existing Karl Storz endoscopes, light sources, and accessories.</p> <p>The upgraded system shall be <b>fully backward compatible with existing Karl Storz rigid and flexible video endoscope</b> which which was previously utilized with the following system detail:</p> <table border="1"> <tr> <td colspan="3"><b>Brand: Karl Storz</b></td> </tr> <tr> <td></td> <td><b>Camera Control Unit</b></td> <td><b>Light Source</b></td> </tr> <tr> <td><b>Unit 1</b></td> <td><b>Image 1 Hub</b></td> <td><b>Xenon 300</b></td> </tr> <tr> <td><b>Unit 2</b></td> <td><b>Image 1 Hub</b></td> <td><b>Xenon 300</b></td> </tr> </table>		<b>Brand: Karl Storz</b>				<b>Camera Control Unit</b>	<b>Light Source</b>	<b>Unit 1</b>	<b>Image 1 Hub</b>	<b>Xenon 300</b>	<b>Unit 2</b>	<b>Image 1 Hub</b>	<b>Xenon 300</b>
<b>Brand: Karl Storz</b>														
	<b>Camera Control Unit</b>	<b>Light Source</b>												
<b>Unit 1</b>	<b>Image 1 Hub</b>	<b>Xenon 300</b>												
<b>Unit 2</b>	<b>Image 1 Hub</b>	<b>Xenon 300</b>												
1.2	The system shall be designed for use in an <b>operating theatre environment</b> and suitable for continuous clinical operation													
1.3	<b>SYSTEM CONFIGURATION</b> Each camera system upgrade shall consist of, but not limited to: <ol style="list-style-type: none"> <li>1. Autoclavable camera head</li> <li>2. Camera Control Unit (CCU)</li> </ol>													
1.4	<b>CAMERA HEAD</b>													
1.4.1	High performance <b>CMOS camera head</b>													
1.4.2	Native resolution: <b>Full HD (1920 × 1080) or higher</b>													
1.4.3	Frame rate: 50/60 Hz or equivalent													
1.4.4	Ergonomic, lightweight design suitable for prolonged operative procedures													
1.4.5	Programmable camera head buttons for: <ul style="list-style-type: none"> <li>▪ Image capture</li> <li>▪ Video recording</li> <li>▪ Digital zoom</li> <li>▪ White balance</li> </ul>													
1.4.6	Camera head shall be: <ul style="list-style-type: none"> <li>▪ Soakable</li> <li>▪ Sterilizable with Hydrogen Peroxide</li> <li>▪ Disinfected with High-Level Disinfection</li> </ul>													
1.4.7	Fully compatible with the offered camera control unit													
1.5	<b>CAMERA CONTROL UNIT</b>													
1.5.1	Advanced image processing unit must support at least Full HD or better													
1.5.2	The system shall be for use with camera head, rigid and flexible endoscopes for laparoscopic and gynaecological visualization													
1.5.3	Real-time image enhancement technologies including: <ul style="list-style-type: none"> <li>▪ Brightness optimization</li> <li>▪ Color contrast enhancement</li> <li>▪ Tissue structure and vascular enhancement modes</li> </ul>													
1.5.4	Automatic and manual white balance													
1.5.5	Shall support multiple simultaneous video output, including but not limited to: <ul style="list-style-type: none"> <li>▪ HDMI (Tenderer to specify HDMI Output quantity)</li> <li>▪ DVI-D (Tenderer to specify DVI-D Output quantity)</li> <li>▪ SDI (Tenderer to specify SDI Output quantity)</li> </ul>													
1.5.6	Low-latency video signal transmission suitable for real-time surgical procedures													
1.5.7	USB interface available for Data export and software updates													
1.5.8	Shall allow automatic device recognition and coordinated control to improve workflow efficiency													
1.5.9	Shall enable interconnection and synchronized operation of: <ul style="list-style-type: none"> <li>▪ Camera control units</li> <li>▪ Light sources</li> </ul>													

<b>SECTION 1 – USER REQUIREMENTS</b>	
<b>REF. NO.</b>	<b>DESCRIPTION</b>
	<ul style="list-style-type: none"> <li>▪ Insufflators</li> <li>▪ Documentation systems</li> </ul>
1.6	Tenderer may offer alternative brand(s) of system, provided that the scope is fully compatible with the existing rigid and flexible videoscope, including the provision of all necessary compatible connectors, at no additional cost.
1.7	<b>SAFETY STANDARD REQUIREMENT</b>
1.7.1	Compliance with the following standard or equivalent: <ul style="list-style-type: none"> <li>▪ IEC 60601-1 (Medical Electrical Equipment)</li> <li>▪ IEC 60601-1-2 (Electromagnetic Compatibility)</li> <li>▪ CE and/or FDA approval</li> </ul>
<b>2</b>	<b>CONSUMABLES AND ACCESSORIES</b>
2.1	Inclusive of all the accessories for the machine to be fully functional
2.2	Each camera head system shall be supplied with a camera head container suitable for sterilization and storage
	* In your quotation/tender document, please breakdown/itemized the price for each accessories/ consumables
<b>3</b>	<b>END USER AND TECHNICAL TRAINING</b>
3.1	Conduct <b>user training</b> to the all-end users by an application specialist or competent local engineer including but not limited to: <ul style="list-style-type: none"> <li>▪ Basic user operation, user troubleshooting and user maintenance</li> <li>▪ Provide Operating manual (Hardcopy and/or Softcopy)</li> </ul>
3.1.1	Tenderer must <b>prepare a training attendance or proof of training done to end user during commissioning and the refresher course (6) months after commissioning.</b>
3.2	<b>Introductory Technical Training</b> to Biomedical Engineers and Technicians at BME Office by competent Tenderer's Engineer/Technicians that includes but not limited to: <ul style="list-style-type: none"> <li>▪ Troubleshooting and basic corrective maintenance</li> <li>▪ Handling and basic inspection maintenance</li> </ul> *(Two sessions/groups if required)
<b>4</b>	<b>WARRANTY</b>
4.1	Tenderer to include warranty period of <b>at least TWO (2) years</b>
4.2	Tenderers to <b>ACKNOWLEDGE</b> 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: <ul style="list-style-type: none"> <li>▪ Scope of Warranty</li> <li>▪ One-time Planned Preventive Maintenance Per Year during warranty</li> <li>▪ Comprehensive Corrective Maintenance of Main Unit</li> </ul>
<b>5</b>	<b>PRODUCT DEMONSTRATION (UPON REQUIREMENT)</b>
5.1	If requested by the Government, vendor shall provide a product demonstration as part of the evaluation process The tenderer shall provide either of the following demonstration modes: <ul style="list-style-type: none"> <li>▪ Physical Demonstration (In-Person): A complete working unit must be brought to the designated demonstration site as specified by the procuring entity. The demonstration shall be conducted by qualified personnel from the supplier.</li> <li>▪ Online Demonstration (Virtual): The demonstration may be conducted via a live video conferencing platform (e.g., Zoom, MS Teams). The session must include: <ul style="list-style-type: none"> <li>○ Live operation of the actual product.</li> <li>○ Real-time interaction to address questions or perform requested functions.</li> <li>○ High-definition video and clear audio for full visibility and understanding.</li> </ul> </li> </ul>

\* In your quotation/tender document, please breakdown/itemized the price for each accessories/ consumables

<b>SECTION 2 – PRICE PROPOSAL</b>	
<b>PURCHASE PRICE</b>	<b>PER UNIT</b>
	<b>TOTAL</b>

<b>SECTION 3 - PROCUREMENT AND TECHNICAL SPECIFICATION</b>	
<b>BRAND:</b>	<b>MODEL:</b>
<b>COUNTRY OF ORIGIN:</b>	<b>YEAR INTRODUCED TO MARKET:</b>
<b>WARRANTY PERIOD:</b>	<b>LAST COUNTRY SOLD TO:</b>
<b>PRICE VALIDITY: [AT LEAST <u>ONE (1) YEAR</u> PRICE VALIDTY]</b>	<b>DELIVERY TIME:</b>

<b>SECTION 3 - PROCUREMENT AND TECHNICAL SPECIFICATION</b>	
<b>AUTHORIZED DISTRIBUTOR: (AUTHORIZED DISTRIBUTOR LETTER ATTACHED)</b>	
<b>DETAILED BROCHURE INCLUDED</b>	
<b>USER AND SERVICE MANUALS:</b>	
<b>MAINS POWER SUPPLY:</b>	
<b>BATTERY</b>	
<b>POWER ADAPTER/CHARGER OUTPUT RATING:</b>	
<b>EQUIPMENT AMBIENT OPERATING TEMPERATURE RANGE:</b>	
<b>INTERNATIONAL SAFETY STANDARD</b> Must comply to at least 1 safety Standards and certification (Please attached the copy of stated standards and certifications)	
<b>NUMBER OF TECHNICAL SUPPORT (ENGINEER/TECHNICIAN)</b> Please provide training or certification for locals who is trained/certified	<b>LOCAL</b>
	<b>OVERSEA (SPECIFY LOCATION)</b>
<b>DIMENSIONS AND WEIGHT OF MAIN UNIT:</b>	
<b>EQUIPMENT WHOLE LIFETIME SUPPORT:</b>	

## 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:
  - 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.
- **Two time Planned Preventive Maintenance (PPM) PER YEAR** 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 below to include items or terms which is not listed in the exclusion list above for MOH consideration.

NO.	TERMS AND CONDITIONS
1	Tenderer must be registered with the Ministry of Health.
2	<b>TENDER FORM should be filled</b> completely including the <b>USER REQUIREMENT FORM</b> (if available). Submission of incomplete form <b>MAY</b> cause <b>DISQUALIFICATION OF TENDER</b> .
3	Each tenderer is allowed to quote <b>ONE BRAND WITH ONE PRICE ONLY</b> for each item. Submission of more than one brand and price will cause <b>DISQUALIFICATION OF TENDER</b> .
4	All consumables supplied throughout this tender <u>shall</u> have a minimum expiry date of <b>twelve (12) months / on delivery</b> (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).
5	Brochures / catalogues should be submitted / attached with tender document.
6	Any room renovation which may be required, it is mandatory to conduct site visit (if applicable)
7	Samples should be submitted together with tender or within fourteen (14 days) of the tender closing dates (if applicable).
8	<b>DELIVERY PERIOD:</b> (Please state) Not More Than <b>90 days</b> upon confirmation
9	<b>PRICE VALIDITY:</b> The quotation shall remain valid for <b>12 MONTHS</b> 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).
10	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. During <b>delivery</b> , the vendor is required to <b>provide proof of manufacture date confirming the equipment is new</b> .

**SECTION 3  
TENDER FORM**

To:

**TENDER REFERENCE NO: KK/101/2026/HTD**

**INVITATION TO TENDER  
SUPPLY, DELIVERY, INSTALLATION, TESTING, COMMISSIONING AND UPGRADE OF EXISTING  
SURGICAL ENDOSCOPY CAMERA HEAD SYSTEM FOR OPERATING THEATRE, RIPAS  
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<b>DELIVERY PERIOD</b>	
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<b>SCOPE OF WORK AND SUMMARY OF PRICES</b>
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**This tender is for the Supply, Delivery, Installation, Testing, Commissioning, Warranty and  
Maintenance of:**

<b>DESCRIPTION</b>	<b>QUANTITY</b>	<b>YES</b>	<b>NO</b>	<b>REMARKS</b>
<b>UPGRADE OF EXISTING SURGICAL ENDOSCOPY CAMERA HEAD SYSTEM</b>	<b>2</b>			

SECTION 1 – USER REQUIREMENTS																
REF. NO.	DESCRIPTION	(✓)		STATE OR SPECIFY OR REMARKS OR BROCHURE PAGE												
		Y	N													
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SECTION 1 – USER REQUIREMENTS				
REF. NO.	DESCRIPTION	(✓)		STATE OR SPECIFY OR REMARKS OR BROCHURE PAGE
		Y	N	
	<ul style="list-style-type: none"> <li>▪ SDI (<a href="#">Tenderer to specify SDI Output quantity</a>)</li> </ul>			
1.5.6	Low-latency video signal transmission suitable for real-time surgical procedures			
1.5.7	USB interface available for Data export and software updates			
1.5.8	Shall allow automatic device recognition and coordinated control to improve workflow efficiency			
1.5.9	Shall enable interconnection and synchronized operation of: <ul style="list-style-type: none"> <li>▪ Camera control units</li> <li>▪ Light sources</li> <li>▪ Insufflators</li> <li>▪ Documentation systems</li> </ul>			
1.6	Tenderer may offer alternative brand(s) of system, provided that the scope is fully compatible with the existing rigid and flexible videoscope, including the provision of all necessary compatible connectors, at no additional cost.			
1.7	<b>SAFETY STANDARD REQUIREMENT</b>			
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2	<b>CONSUMABLES AND ACCESSORIES</b>			
2.1	Inclusive of all the accessories for the machine to be fully functional			
2.2	Each camera head system shall be supplied with a camera head container suitable for sterilization and storage			
	<i>* In your quotation/tender document, please breakdown/itemized the price for each accessories/ consumables</i>			
3	<b>END USER AND TECHNICAL TRAINING</b>			
3.1	Conduct <b>user training</b> to the all-end users by an application specialist or competent local engineer including but not limited to: <ul style="list-style-type: none"> <li>▪ Basic user operation, user troubleshooting and user maintenance</li> <li>▪ Provide Operating manual (Hardcopy and/or Softcopy)</li> </ul>			
3.1.1	Tenderer must <b>prepare a training attendance or proof of training done to end user during commissioning and the refresher course (6) months after commissioning.</b>			
3.2	<b>Introductory Technical Training</b> to Biomedical Engineers and Technicians at BME Office by competent Tenderer's Engineer/Technicians that includes but not limited to: <ul style="list-style-type: none"> <li>▪ Troubleshooting and basic corrective maintenance</li> <li>▪ Handling and basic inspection maintenance</li> </ul> <i>*(Two sessions/groups if required)</i>			
4	<b>WARRANTY</b>			
4.1	Tenderer to include warranty period of <b>at least TWO (2) years</b>			
4.2	Tenderers to <b>ACKNOWLEDGE</b> 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:			

SECTION 1 – USER REQUIREMENTS				
REF. NO.	DESCRIPTION	(✓)		STATE OR SPECIFY OR REMARKS OR BROCHURE PAGE
		Y	N	
	<ul style="list-style-type: none"> <li>▪ Scope of Warranty</li> <li>▪ One-time Planned Preventive Maintenance Per Year during warranty</li> <li>▪ Comprehensive Corrective Maintenance of Main Unit</li> </ul>			
5	<b>PRODUCT DEMONSTRATION (UPON REQUIREMENT)</b>			
5.1	<p>If requested by the Government, vendor shall provide a product demonstration as part of the evaluation process The tenderer shall provide either of the following demonstration modes:</p> <ul style="list-style-type: none"> <li>▪ Physical Demonstration (In-Person): A complete working unit must be brought to the designated demonstration site as specified by the procuring entity. The demonstration shall be conducted by qualified personnel from the supplier.</li> <li>▪ Online Demonstration (Virtual): The demonstration may be conducted via a live video conferencing platform (e.g., Zoom, MS Teams). The session must include: <ul style="list-style-type: none"> <li>○ Live operation of the actual product.</li> <li>○ Real-time interaction to address questions or perform requested functions.</li> <li>○ High-definition video and clear audio for full visibility and understanding.</li> </ul> </li> </ul>			

SECTION 2 – PRICE PROPOSAL		
PURCHASE PRICE	PER UNIT	BND\$
	TOTAL	BND\$

SECTION 3 - PROCUREMENT AND TECHNICAL SPECIFICATION			
BRAND:		MODEL:	
COUNTRY OF ORIGIN:		YEAR INTRODUCED TO MARKET:	
WARRANTY PERIOD:		LAST COUNTRY SOLD TO:	
PRICE VALIDITY: [AT LEAST <u>ONE (1) YEAR</u> PRICE VALIDTY]		DELIVERY TIME:	

SECTION 3 - PROCUREMENT AND TECHNICAL SPECIFICATION				
<b>AUTHORIZED DISTRIBUTOR: (AUTHORIZED DISTRIBUTOR LETTER ATTACHED)</b>	APPOINTED BRUNEI DISTRIBUTOR			
	PROCURE FROM OVERSEA AUTHORIZED DISTRIBUTOR	COMPANY NAME:		
<b>DETAILED BROCHURE INCLUDED</b>	YES		NO	<input checked="" type="checkbox"/> or specify where appropriate
<b>USER AND SERVICE MANUALS:</b>	YES		NO	Tenderers to acknowledge that they must provide at least <b>TWO</b> sets of <b>USER AND SERVICE</b> manuals when applying commissioning form. One Set for End User, One Set for BME. (Please provide hardcopy or softcopy)
<b>MAINS POWER SUPPLY:</b>	220V-240V		OTHERS:	
	50-60HZ		OTHERS:	
<b>BATTERY</b>	RECHARGEABLE		SINGLE-USE	REPLACEABLE
	OTHERS:			
	TYPE OF BATTERY:			
	RATING:			
<b>POWER ADAPTER/CHARGER OUTPUT RATING:</b>				
<b>EQUIPMENT AMBIENT OPERATING TEMPERATURE RANGE:</b>				
<b>INTERNATIONAL SAFETY STANDARD</b> Must comply to at least 1 safety Standards and certification (Please attached the copy of stated standards and certifications)			<input checked="" type="checkbox"/> Tick where appropriate <input type="checkbox"/> US FDA Standard, <input type="checkbox"/> European Union CE MARK, <input type="checkbox"/> Australian TGA Standard, <input type="checkbox"/> Canadian CSA Standard or <input type="checkbox"/> Japanese JIS Standard. Others (Please specify): _____	
<b>NUMBER OF TECHNICAL SUPPORT (ENGINEER/TECHNICIAN)</b>  Please provide training or certification for locals who is trained/certified	LOCAL		<input type="checkbox"/> Trained / Certified <input type="checkbox"/> Not yet trained on the product	
	OVERSEA (SPECIFY LOCATION)		NEAREST LOCATION:	
<b>DIMENSIONS AND WEIGHT OF MAIN UNIT:</b>		<input type="checkbox"/> mm <input type="checkbox"/> cm <input type="checkbox"/> inch		<input type="checkbox"/> Kilogram (Kg) <input type="checkbox"/> Gram(g) <input type="checkbox"/> Pound (lbs)
<b>EQUIPMENT WHOLE LIFE TIME SUPPORT:</b>	The supplier shall ensure that spare parts for the equipment are available for a minimum of 10 years after installation, with the support period extending beyond the expected lifecycle of the equipment. No of years: _____ (Please specify)			

**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:
  - 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
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  - C. If a unit or accessory is deemed used item or refurbished item (not a new unit) by the user and BME Unit.
- **Two time Planned Preventive Maintenance (PPM) PER YEAR** 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 below to include items or terms which is not listed in the exclusion list above for MOH consideration.

**TENDERER ACKNOWLEDGMENT**

**COMPANY CHOP AND SIGNATURE**

NO.	TERMS AND CONDITIONS	VENDOR'S OFFER (PLEASE STATE)
1	Tenderer must be registered with the Ministry of Health.	
2	<b>TENDER FORM should be filled</b> completely including the <b>USER REQUIREMENT FORM</b> (if available). Submission of incomplete form <b>MAY</b> cause <b>DISQUALIFICATION OF TENDER</b> .	
3	Each tenderer is allowed to quote <b>ONE BRAND WITH ONE PRICE ONLY</b> for each item. Submission of more than one brand and price will cause <b>DISQUALIFICATION OF TENDER</b> .	
4	All consumables supplied throughout this tender <u>shall</u> have a minimum expiry date of <b>twelve (12) months / on delivery</b> (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).	
5	Brochures / catalogues should be submitted / attached with tender document.	
6	Any room renovation which may be required, it is mandatory to conduct site visit (if applicable)	
7	Samples should be submitted together with tender or within fourteen (14 days) of the tender closing dates (if applicable).	
8	<b>DELIVERY PERIOD:</b> (Please state) Not More Than <b>90 days</b> upon confirmation	
9	<b>PRICE VALIDITY:</b> The quotation shall remain valid for <b>12 MONTHS</b> 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).	
10	The equipment supplied must be <b>newly manufactured, unused, and in its original, sealed packaging</b> . The equipment must not be previously owned, refurbished, or reconditioned in any form. During <b>delivery</b> , the vendor is required to <b>provide proof of manufacture date confirming the equipment is new</b> .	