

TENDER REF. NO.: KK/202/2026/UPP

**MINISTRY OF HEALTH
NEGARA BRUNEI DARUSSALAM**

**THE SUPPLY, DELIVERY, INSTALLATION, TESTING AND
COMMISSIONING OF WASHER DISINFECTOR FOR
CENTRAL STERILE SERVICES DEPARTMENT (CSSD),
RAJA ISTERI PENGIRAN ANAK SALEHA (RIPAS)
HOSPITAL**

TENDER FEE : \$30.00

RECEIPT NO. :

CLOSING DATE : ON TUESDAY, 06th October 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

To:

TENDER REFERENCE NO: KK/202/2026/UPP

INVITATION TO TENDER
SUPPLY, DELIVERY, INSTALLATION, TESTING AND COMMISSIONING OF WASHER
DISINFECTOR FOR CENTRAL STERILE SERVICES DEPARTMENT (CSSD), RAJA ISTERI
PENGIRAN ANAK SALEHA (RIPAS) HOSPITAL

TENDER OF (*name of tenderer*) : _____

Company/Business Registration No. : _____

Tender Closing Date : _____

SCOPE OF WORK	
Item 1 - Supply of <u>1 unit</u> of Freestanding washer disinfecter for general use in CSSD, RIPAS HOSPITAL	
Item 2 - Supply of <u>1 unit</u> of Freestanding washer disinfecter for Milk Bottles in CSSD, RIPAS HOSPITAL	

	SECTION 1 – USER REQUIREMENTS
ITEM 1	WASHER DISINFECTOR – FOR GENERAL USE
1.1	Medical-grade washer disinfecter suitable for CSSD use
1.2	Suitable for cleaning and thermal disinfection of reusable surgical instruments, containers, anaesthesia accessories, MIS instruments, and medical devices.
1.3	Configuration: Freestanding, single-door pass-through type suitable for barrier installation between dirty and clean areas
1.4	Construction
1.4.1	Chamber and exterior constructed from stainless steel
1.4.2	Automatic or electrically assisted double-door system with glass viewing panel or equivalent.
1.4.3	Illuminated washing chamber.
1.4.4	Compact footprint suitable for CSSD installation space
1.4.5	Front access for maintenance and servicing.
1.4.6	Chamber floor designed for effective drainage.

	SECTION 1 – USER REQUIREMENTS
1.4.7	Complete with integrated lower cabinet/compartment for storage of chemical containers
1.5	Washing Chamber and Capacity
1.5.1	Chamber capacity: At least 350 litres or better
1.5.2	Suitable for approximately 12–18 DIN trays or equivalent
1.5.3	Suitable for processing surgical instruments, MIS instruments, anaesthesia accessories, containers, and other reusable medical devices.
1.6	Washing and disinfection system
1.6.1	Fully automatic PLC or microprocessor-controlled operation.
1.6.2	Equipped with rotating wash arms at top and bottom of chamber, including rack-level wash arms where applicable.
1.6.3	Thermal disinfection system compliant with EN ISO 15883.
1.6.4	Comes with at least 3 dosing pumps for detergents and disinfectants
1.6.5	Wash/disinfection temperature shall be monitored by dual independent temperature sensors for process validation and redundancy
1.6.6	Automatic self-disinfection program for chamber and piping system.
1.6.7	Integrated drying system with HEPA-filtered air.
1.6.8	Built-in vapor condenser or equivalent system to minimise steam release into the room.
1.6.9	Smart water management or equivalent system for reduced water and chemical consumption.
1.7	Control System
1.7.1	Colour touchscreen or LCD control panel.
1.7.2	Standard programmable washing and disinfection cycles available.
1.7.3	Capable of storing, downloading, and reviewing cycle data
1.7.4	USB and/or network connectivity for documentation and traceability purposes.
1.7.5	Comes with printer
1.8	Safety Features

	SECTION 1 – USER REQUIREMENTS
1.8.1	Visual and audible alarm system.
1.8.2	Emergency stop button.
1.8.3	Automatic door lock after program start
1.8.4	Backflow prevention system.
1.8.5	Independent temperature monitoring system.
1.8.6	Door interlock system to prevent simultaneous opening of doors
1.9	Safety Standard Requirement Compliance with the following standard or equivalent: ▪ IEC and/or EN Safety Standards ▪ CE and/or FDA approval
1.10	Accessories/consumables to be supplied with each unit: Inclusive of all the accessories for the machine to be fully functional, including but not limited to:
1.10.1	Two (2) units of Transport trolley
1.10.2	One (1) set of 6-Level rack to accommodate 18 DIN instrument trays or equivalent
1.10.3	One (1) set of Basic rack suitable for utensils, bowls, dishes & glassware
1.10.4	One (1) set of Anaesthesia rack with tube tower with capacity of 21 breathing hose or equivalent
1.10.5	Eighteen (18) sets of Basket moulded DIN trays for surgical instruments
1.10.6	One (1) set of large size mesh basket
1.10.7	One (1) set of Insert for trays/tablets for loading
1.10.8	One (1) set of Tray for Small Anaesthesia Instrument Parts
1.10.9	One (1) set of Insert for Kidney Dishes
1.10.10	One (1) set of Insert for Universal insert with pins for fixing glasses and utensils
ITEM 2	WASHER DISINFECTOR FOR MILK BOTTLES
2.1	Medical-grade washer disinfector suitable for cleaning and thermal disinfection of reusable milk bottles, feeding bottles, feeding accessories, and related neonatal feeding equipment
2.2	Configuration: Freestanding, single-door pass-through type suitable for barrier installation between dirty and clean areas
2.3	Construction

	SECTION 1 – USER REQUIREMENTS
2.3.1	Chamber and exterior constructed from stainless steel.
2.3.2	Glass viewing panel or equivalent door design.
2.3.3	Illuminated washing chamber.
2.3.4	Compact footprint suitable for designated installation space.
2.3.5	Front access for maintenance and servicing
2.3.6	Chamber floor designed for effective drainage
2.4	Washing Chamber and Capacity
2.4.1	Chamber capacity: At least 200 litres or better
2.4.2	Suitable for processing milk bottles, teats, feeding containers, breast pump accessories, and related neonatal feeding accessories.
2.4.3	Rack system shall support secure positioning of bottles and accessories during washing and disinfection process.
2.5	Washing and Disinfection System
2.5.1	Fully automatic PLC or microprocessor-controlled operation
2.5.2	Equipped with rotating wash arms and injection washing system where applicable.
2.5.3	Thermal disinfection system compliant with EN ISO 15883.
2.5.4	Automatic self-disinfection program for chamber and piping system.
2.5.5	Integrated drying system with HEPA-filtered air.
2.5.6	Built-in vapor condenser or equivalent system to minimise steam release into the room.
2.5.7	Adjustable washing and drying parameters.
2.6	Control System
2.6.1	Colour touchscreen or LCD control panel.
2.6.2	Standard programmable washing and disinfection cycles available
2.6.3	Capable of storing, downloading, and reviewing cycle data.

	SECTION 1 – USER REQUIREMENTS
2.6.4	USB and/or network connectivity for documentation and traceability purposes.
2.6.5	Comes with printer
2.7	Safety Features
2.7.1	Visual and audible alarm system.
2.7.2	Emergency stop button.
2.7.3	Backflow prevention system.
2.7.4	Independent temperature monitoring system.
2.7.5	Door safety interlock system
2.8	Safety Standard Requirement Compliance with the following standard or equivalent: ▪ IEC and/or EN Safety Standards ▪ CE and/or FDA approval
2.9	Accessories/consumables to be supplied with each unit: Inclusive of all the accessories for the machine to be fully functional, including but not limited to:
2.9.1	Two (2) units of Transport trolley
2.9.2	Two (2) units of Milk bottle washing rack
2.9.3	Basket for feeding accessories

SECTION 3

TENDER FORM

To:

TENDER REFERENCE NO: KK/202/2026/UPP

INVITATION TO TENDER
SUPPLY, DELIVERY, INSTALLATION, TESTING AND COMMISSIONING OF WASHER DISINFECTOR FOR CENTRAL STERILE SERVICES
DEPARTMENT (CSSD), RAJA ISTERI PENGIRAN ANAK SALEHA (RIPAS) HOSPITAL

TENDER OF (*name of tenderer*) : _____

Company/Business Registration No. : _____

Tender Closing Date : _____

SCOPE OF WORK			
Please ☒ Tick where appropriate	Yes	No	Remarks
Item 1 - Supply of <u>1 unit</u> of Freestanding washer disinfector for general use in CSSD, RIPAS HOSPITAL			
Item 2 - Supply of <u>1 unit</u> of Freestanding washer disinfector for Milk Bottles in CSSD, RIPAS HOSPITAL			

SECTION 1 – USER REQUIREMENTS				
Please ☒ Tick where appropriate		Yes	No	Remarks
ITEM 1	WASHER DISINFECTOR – FOR GENERAL USE			
1.1	Medical-grade washer disinfector suitable for CSSD use			
1.2	Suitable for cleaning and thermal disinfection of reusable surgical instruments, containers, anaesthesia accessories, MIS instruments, and medical devices.			
1.3	Configuration: Freestanding, single-door pass-through type suitable for barrier installation between dirty and clean areas			
1.4	Construction			
1.4.1	Chamber and exterior constructed from stainless steel			
1.4.2	Automatic or electrically assisted double-door system with glass viewing panel or equivalent.			
1.4.3	Illuminated washing chamber.			
1.4.4	Compact footprint suitable for CSSD installation space			
1.4.5	Front access for maintenance and servicing.			
1.4.6	Chamber floor designed for effective drainage.			
1.4.7	Complete with integrated lower cabinet/compartment for storage of chemical containers			
1.5	Washing Chamber and Capacity			

SECTION 1 – USER REQUIREMENTS				
Please ☒ Tick where appropriate		Yes	No	Remarks
1.5.1	Chamber capacity: At least 350 litres or better			
1.5.2	Suitable for approximately 12–18 DIN trays or equivalent			
1.5.3	Suitable for processing surgical instruments, MIS instruments, anaesthesia accessories, containers, and other reusable medical devices.			
1.6	Washing and disinfection system			
1.6.1	Fully automatic PLC or microprocessor-controlled operation.			
1.6.2	Equipped with rotating wash arms at top and bottom of chamber, including rack-level wash arms where applicable.			
1.6.3	Thermal disinfection system compliant with EN ISO 15883.			
1.6.4	Comes with at least 3 dosing pumps for detergents and disinfectants			
1.6.5	Wash/disinfection temperature shall be monitored by dual independent temperature sensors for process validation and redundancy			
1.6.6	Automatic self-disinfection program for chamber and piping system.			
1.6.7	Integrated drying system with HEPA-filtered air.			
1.6.8	Built-in vapor condenser or equivalent system to minimise steam release into the room.			
1.6.9	Smart water management or equivalent system for reduced water and chemical consumption.			

SECTION 1 – USER REQUIREMENTS				
Please ☒ Tick where appropriate		Yes	No	Remarks
1.7	Control System			
1.7.1	Colour touchscreen or LCD control panel.			
1.7.2	Standard programmable washing and disinfection cycles available.			
1.7.3	Capable of storing, downloading, and reviewing cycle data			
1.7.4	USB and/or network connectivity for documentation and traceability purposes.			
1.7.5	Comes with printer			
1.8	Safety Features			
1.8.1	Visual and audible alarm system.			
1.8.2	Emergency stop button.			
1.8.3	Automatic door lock after program start			
1.8.4	Backflow prevention system.			
1.8.5	Independent temperature monitoring system.			
1.8.6	Door interlock system to prevent simultaneous opening of doors			

SECTION 1 – USER REQUIREMENTS				
Please ☒ Tick where appropriate		Yes	No	Remarks
1.9	Safety Standard Requirement Compliance with the following standard or equivalent: ▪ IEC and/or EN Safety Standards ▪ CE and/or FDA approval			
1.10	Accessories/consumables to be supplied with each unit: Inclusive of all the accessories for the machine to be fully functional, including but not limited to:			
1.10.1	Two (2) units of Transport trolley			
1.10.2	One (1) set of 6-Level rack to accommodate 18 DIN instrument trays or equivalent			
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1.10.4	One (1) set of Anaesthesia rack with tube tower with capacity of 21 breathing hose or equivalent			
1.10.5	Eighteen (18) sets of Basket moulded DIN trays for surgical instruments			
1.10.6	One (1) set of large size mesh basket			
1.10.7	One (1) set of Insert for trays/tablets for loading			
1.10.8	One (1) set of Tray for Small Anaesthesia Instrument Parts			
1.10.9	One (1) set of Insert for Kidney Dishes			
1.10.10	One (1) set of Insert for Universal insert with pins for fixing glasses and utensils			

SECTION 1 – USER REQUIREMENTS				
Please ☒ Tick where appropriate		Yes	No	Remarks
ITEM 2	WASHER DISINFECTOR FOR MILK BOTTLES			
2.1	Medical-grade washer disinfector suitable for cleaning and thermal disinfection of reusable milk bottles, feeding bottles, feeding accessories, and related neonatal feeding equipment			
2.2	Configuration: Freestanding, single-door pass-through type suitable for barrier installation between dirty and clean areas			
2.3	Construction			
2.3.1	Chamber and exterior constructed from stainless steel.			
2.3.2	Glass viewing panel or equivalent door design.			
2.3.3	Illuminated washing chamber.			
2.3.4	Compact footprint suitable for designated installation space.			
2.3.5	Front access for maintenance and servicing			
2.3.6	Chamber floor designed for effective drainage			
2.4	Washing Chamber and Capacity			
2.4.1	Chamber capacity: At least 200 litres or better			
2.4.2	Suitable for processing milk bottles, teats, feeding containers, breast pump accessories, and related neonatal feeding accessories.			

SECTION 1 – USER REQUIREMENTS				
Please ☒ Tick where appropriate		Yes	No	Remarks
2.4.3	Rack system shall support secure positioning of bottles and accessories during washing and disinfection process.			
2.5	Washing and Disinfection System			
2.5.1	Fully automatic PLC or microprocessor-controlled operation			
2.5.2	Equipped with rotating wash arms and injection washing system where applicable.			
2.5.3	Thermal disinfection system compliant with EN ISO 15883.			
2.5.4	Automatic self-disinfection program for chamber and piping system.			
2.5.5	Integrated drying system with HEPA-filtered air.			
2.5.6	Built-in vapor condenser or equivalent system to minimise steam release into the room.			
2.5.7	Adjustable washing and drying parameters.			
2.6	Control System			
2.6.1	Colour touchscreen or LCD control panel.			
2.6.2	Standard programmable washing and disinfection cycles available			
2.6.3	Capable of storing, downloading, and reviewing cycle data.			

SECTION 1 – USER REQUIREMENTS				
Please ☒ Tick where appropriate		Yes	No	Remarks
2.6.4	USB and/or network connectivity for documentation and traceability purposes.			
2.6.5	Comes with printer			
2.7	Safety Features			
2.7.1	Visual and audible alarm system.			
2.7.2	Emergency stop button.			
2.7.3	Backflow prevention system.			
2.7.4	Independent temperature monitoring system.			
2.7.5	Door safety interlock system			
2.8	Safety Standard Requirement Compliance with the following standard or equivalent: ▪ IEC and/or EN Safety Standards ▪ CE and/or FDA approval			
2.9	Accessories/consumables to be supplied with each unit: Inclusive of all the accessories for the machine to be fully functional, including but not limited to:			
2.9.1	Two (2) units of Transport trolley			
2.9.2	Two (2) units of Milk bottle washing rack			

SECTION 1 – USER REQUIREMENTS				
Please ☒ Tick where appropriate		Yes	No	Remarks
2.9.3	Basket for feeding accessories			

Please ☒ Tick where appropriate		Yes	No	Remarks
3	INSTALLATION REQUIREMENT			
3.1	Washer disinfectpr shall be compatible with and fit within the existing installation space designated for the washer disinfectpr at CSSD, RIPAS Hospital, including any necessary wall hacking works and renovation required for installation			
3.2	Tenderer shall provide installation service where necessary, including but not limited to the following: ▪ Site preparation and planning works, including new vinyl flooring and antibacterial wall painting ▪ Ensure utilities meet system requirements, including electrical supply, water supply, and drainage connection where applicable. Any additional electrical works, utility upgrades, or connections required for installation and commissioning shall be included. ▪ Provision of installation drawings and layout guidance ▪ System delivery, assembly, installation, and configuration ▪ Support for acceptance testing, commissioning, and rectification of defects.			
3.3	All associated renovation works, including materials, labour, and finishing, shall be included in the Tenderer's offer			
3.4	Tenderer shall be responsible to mobilize and demobilize the machineries used at the work site and to remove the debris from the site after completion works.			
3.5	Mandatory Site Visit A mandatory site visit shall be conducted as part of the tendering process. Tenderers are required to contact the BME personnel in charge via email at Nuramaliah.Jamaludin@moh.gov.bn. The Site Visit Form is attached as Annex A.			
4	END USER AND TECHNICAL TRAINING			

Please ☒ Tick where appropriate		Yes	No	Remarks
4.1	Conduct user training to the all-end users by an application specialist or competent local engineer including but not limited to: ▪ Basic user operation, user troubleshooting and user maintenance ▪ Provide Operating manual (Hardcopy and/or Softcopy)			
4.1.1	Tenderer must prepare a training attendance or proof of training done to end user during commissioning and the refresher course (6) months after commissioning.			
4.2	Introductory Technical Training to Biomedical Engineers and Technicians at BME Office by competent Tenderer's Engineer/Technicians that includes but not limited to: ▪ Troubleshooting and basic corrective maintenance ▪ Handling and basic inspection maintenance *(Two sessions/groups if required)			
5	WARRANTY			
5.2	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: ▪ Scope of Warranty ▪ One-time Planned Preventive Maintenance Per Year during warranty ▪ Comprehensive Corrective Maintenance of Main Unit			

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 VALIDITY]		DELIVERY TIME:	

SECTION 3 - PROCUREMENT AND TECHNICAL SPECIFICATION					
AUTHORIZED DISTRIBUTOR: (AUTHORIZED DISTRIBUTOR LETTER ATTACHED)	APPOINTED BRUNEI DISTRIBUTOR				
		PROCURE FROM OVERSEA AUTHORIZED DISTRIBUTOR		COMPANY NAME:	
				COMPANY ORIGIN:	
DETAILED BROCHURE INCLUDED		YES		NO	☒ or specify 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 End User, One Set for BME. (Please provide hardcopy or softcopy)
MAINS POWER SUPPLY:		400V		OTHERS:	
		50-60HZ		OTHERS:	
BATTERY		RECHARGEABLE			SINGLE-USE
		OTHERS:			
		TYPE OF BATTERY:			
		RATING:			
POWER ADAPTER/CHARGER OUTPUT RATING:					
EQUIPMENT AMBIENT OPERATING TEMPERATURE RANGE:					
INTERNATIONAL SAFETY STANDARD Must comply to at least 1 safety Standards and certification (Please attached the copy of stated standards and certifications)					☒ Tick where appropriate ☐ US FDA Standard, ☐ European Union CE MARK, ☐ Australian TGA Standard, ☐ Canadian CSA Standard or ☐ Japanese JIS Standard. Others (Please specify):
NUMBER OF TECHNICAL SUPPORT (ENGINEER/TECHNICIAN) Please provide training or certification for locals who is trained/certified	LOCAL		☐ Trained / Certified ☐ Not yet trained on the product		
	OVERSEA (SPECIFY LOCATION)		NEAREST LOCATION:		
DIMENSIONS AND WEIGHT OF MAIN UNIT:		☐ mm ☐ cm ☐ inch			☐ Kilogram (Kg) ☐ Gram(g) ☐ Pound (lbs)
EQUIPMENT WHOLE LIFE TIME SUPPORT:	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 extent 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.

TENDERER ACKNOWLEDGMENT

COMPANY CHOP AND SIGNATURE

ANNEX A: SITE VISIT FORM

TENDER REFERENCE	:	KK/201/2026/UPP	
TENDER TITLE	:	SUPPLY, DELIVERY, INSTALLATION, TESTING AND COMMISSIONING OF WASHER DISINFECTOR FOR CENTRAL STERILE SERVICES DEPARTMENT (CSSD), RAJA ISTERI PENGIRAN ANAK SALEHA (RIPAS) HOSPITAL	
COMPANY NAME	:		COMPANY STAMP AND SIGNATURE
COMPANY FOCAL PERSON CONTACT:	:	NAME: CONTACT NO:	
I hereby on behalf of My Company has made a Site Visit to the following site/location on the date stated below and understand the work requirement(s) and all specification stated in this tender document. I (My Company) also agree not to make any additional claim to MOH should any accident(s) or damage(s) occur during the implementation period.			
<u>FOR OFFICIAL USE ONLY:</u>			
SITE/LOCATION	:	CSSD, RIPASH	VERIFIED BY (BME RIPASH OR CSSD PERSONNEL): NAME: SIGNATURE:
DATE OF SITE VISIT	:		

The Contractor shall visit the site prior to submitting any quotation for the works stated in this Tender and shall satisfy themselves as to the nature of the work and site conditions. The Contractor is required to complete this form and obtain verification signature from the Biomedical Engineer (BME) or End User as proof of the site visit. Failure to comply shall result in disqualification from this Tender

NO.	TERMS AND CONDITIONS	VENDOR'S OFFER (PLEASE STATE)
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 MAY cause DISQUALIFICATION OF TENDER .	
3	Each tenderer is allowed to quote ONE BRAND WITH ONE PRICE ONLY for each item. Submission of more than one brand and price will cause DISQUALIFICATION OF TENDER .	
4	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).	
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	DELIVERY PERIOD: (Please state) Not More Than 90 days upon confirmation	
9	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).	
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 delivery , the vendor is required to provide proof of manufacture date confirming the equipment is new .	

1. We offer and undertake on your acceptance of our Tender to provide the above-mentioned services in accordance with your Invitation To Tender.
2. Our Tender is fully consistent with and does not contradict or derogate from anything in your Invitation To Tender. We have not qualified or changed any of the provisions of your Invitation To Tender.
3. OUR OFFER IS VALID FOR **TWELVE (12)** CALENDAR MONTHS FROM THE TENDER CLOSING DATE.
4. When requested by you, we shall extend the validity of this offer.
5. We further undertake to give you any further information which you may require.

Dated this _____ day of _____, _____

Signature of authorised officer of Tenderer

Name:

Designation:

Tenderer's official stamp