

TENDER REF. NO.: KK/243/2026/HTD

**MINISTRY OF HEALTH
NEGARA BRUNEI DARUSSALAM**

**SUPPLY, DELIVERY, INSTALLATION, TESTING AND
COMMISSIONING ONE (1) UNIT OF GENETIC ANALYSER
SYSTEM FOR HUMAN IDENTIFICATION (HID) FOR
FORENSIC BIOLOGY SECTION, DEPARTMENT OF
SCIENTIFIC SERVICES, MINISTRY OF HEALTH**

TENDER FEE : \$50.00

RECEIPT NO. :

CLOSING DATE : ON TUESDAY, 27th 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 AND REQUIREMENTS

TENDER REFERENCE NO: KK/243/2026/HTD

INVITATION TO TENDER

SUPPLY, DELIVERY, INSTALLATION, TESTING AND COMMISSIONING ONE (1) UNIT OF GENETIC ANALYSER SYSTEM FOR HUMAN IDENTIFICATION (HID) FOR FORENSIC BIOLOGY SECTION, DEPARTMENT OF SCIENTIFIC SERVICES, MINISTRY OF HEALTH

SECTION 1 – USER REQUIREMENTS	
1. STANDARD FEATURES	
<u>NO.</u>	<u>DESCRIPTION</u>
1.0	GENERAL
1.1	Genetic Analyser System for Human Identification (HID) used to amplify forensic samples by polymerase chain reaction and perform fragment analysis by capillary electrophoresis (CE).
1.2	This system shall comprise of the following: • 1 unit of CE instrument. • 2 units of thermal cycler. • Data management and acquisition system. • Accessories. • Performance check and training. • Consumables, reagents and kits for performance check and training. • Documentation of full developmental validation for forensic human identification use.
2	CAPILLARY ELECTROPHORESIS INSTRUMENT SPECIFICATIONS
2.1	Fluorescence-based capillary electrophoresis system.
2.2	Must be able to run on 8-capillary array, upgradeable to 24-capillary when required.
2.3	Allow the use of polymer 4 and polymer 6 for HID (human identification) analysis, fragment analysis and sequencing applications.
2.4	Allow the use of capillary arrays of 36 cm and 50 cm for different fragment analysis and sequencing applications.
2.5	Allow DNA fragment analysis of up to six unique dyes simultaneously.
2.6	Capillary arrays must be able to be used for no less than 160 injections.
2.7	DNA fragment analysis run time of ≤ 40 minutes each.
2.8	Plate capacity of at least two 96-well plates.
2.9	Allow samples to be automatically injected directly from 96-well plates or 8-strip tubes.
2.10	User-friendly consumable installation.
2.11	Consumable tracking with RFID technology.
2.12	Fully automated from reagent loading and replacement, DNA separation, detection and data analysis to generate base-called or size-called results.
2.13	Instrument output files shall be compatible with the GeneMapper ID-x Software v1.6 or equivalent for data processing.
2.14	Long-life, single-line 505nm, solid-state laser excitation source.
2.15	Up to 20kV electrophoresis voltage.
2.16	Active oven temperature control from 18°C to 70°C.
3	THERMAL CYCLER SPECIFICATIONS

3.1	Block format: 96-well, 0.2ml alloy block
3.2	Block range: 6 independent temperature zones (up to 10°C per zone), 30°C range across block
3.3	Maximum block ramp rate: 6.0°C/sec
3.4	Maximum sample ramp rate: 4.4°C/sec
3.5	Temperature accuracy: $\pm 0.25^{\circ}\text{C}$ (35 – 99°C)
3.6	Temperature range: 0 – 100.0°C
3.7	Temperature uniformity: $< 0.5^{\circ}\text{C}$ (30 sec after reaching 95°C)
3.8	Dimensions not more than: Width: 22cm Depth: 25cm Height: 47cm
3.9	PCR volume range: 10 – 100µl
3.10	Instrument memory: USB port and 16GB onboard memory; onboard capacity of > 1000 protocols
3.11	Display interface: Colour digital display
4	DATA MANAGEMENT AND ACQUISITION SYSTEM
4.1	Data management and acquisition system must be supplied for the control of the CE instrument and its auxiliary equipment as well as allowing users for recording and reviewing of the analysis data.
4.2	The system should include: • Two (2) units of stand alone desktop PC with specifications following equipment manufacturer's recommendation. • Communication interface via LAN or WiFi. • Latest processor compatible with the CE instrument data collection software and latest GeneMapper ID-x Software. • One (1) unit of minimum 27" monitor screen and one (1) unit of minimum 34" monitor screen. • Standard keyboard and bluetooth wireless mouse. • Genuine latest Microsoft Windows software compatible to use with the latest GeneMapper ID-x Software. • Two (2) units of high speed external solid state drive with at least 1TB capacity.
4.3	Software Package should include: • Simplified workflow for HID with preconfigured validated protocols and assays for AmpFISTR® kits. • Preliminary data analysis to allow user to evaluate data in real time.
5	ACCESSORIES
5.1	One (1) unit of uninterrupted power supply (UPS) with suitable power supply rating and surge protector (Output voltage uncertainty $\pm 2\%$ or less) must also be provided and connected each to the instrument system and workstation.
5.2	Instrument to be supplied with all necessary standard operating accessories required to ensure successful operation.
5.3	Two (2) units of low-bench working chair.
6	METHOD VALIDATION & TRAINING
6.1	On-site performance check by Human Identification specialists to evaluate the system with Applied Biosystems™ GlobalFiler™ and Yfiler™ Plus Amplification Kits in accordance with the Scientific Working Group DNA Analysis Methods recommendation and/or Quality Assurance Standards for Forensic DNA Testing Laboratories.
6.2	The performance check may include but is not limited to: • Standard Curve and Quality Metrics • Contamination Study

	• Minimum Threshold and Contamination Study • Sensitivity and Stochastic Study • Precision Study: Repeatability and Reproducibility • Accuracy Study: NIST Samples • Known and Non-Probative Sample Study • Mixture Study • Assessment of Non-Allelic Peaks
6.3	On-site training for ALL staff members expected to handle the machine. Please ensure that adequate time is allocated such that training will take place in small groups to minimize staff shortage in the laboratory.
6.4	Certificate of competence is to be issued to all trainees after completion of training.
6.5	The successful tenderer needs to ensure • the key users are updated on the current and/or relevant information related to the system used. • provide ONE (1) off-site professional development in HID DNA analysis for TWO (2) key users. All expenses for attending the training shall be borne by the vendor; full registration, insurance, air ticket, daily allowance, accommodation and transport to and from the airport and training venue.
7	CONSUMABLES, REAGENTS & KITS
7.1	The successful tenderer needs to supply all necessary consumables, reagents and kits required for installation and commissioning, performance check and training.
8	DOCUMENTATION & CERTIFICATION
8.1	Operation manual (1 original and 1 copy).
8.2	Availability of Developmental Validation paper documenting validation studies tested across Applied Biosystems™ STR PCR amplification kits and Precision ID NGS panels. based on requirements listed in the FBI Quality Assurance Standards for Forensic DNA Testing Laboratories and guidelines outlined by the Scientific Working Group on DNA Analysis Methods
8.3	To be supplied during delivery/installation.
9	SITE PREPARATION & ELECTRICAL REQUIREMENTS
9.1	The tenderer shall ensure that the site preparation for the placement of the system taking into the consideration on the safety of the end user during operation of the instrument.
9.2	It is MANDATORY for the tenderer to do site visit prior to tender submission to discuss site requirements. A site visit form to be verified by user during the visit as evidence. Non-attendance will be considered as non-compliance.
9.3	The site preparation details should be listed in the quotation / document submitted.
9.4	All works involving additional electrical supply including wiring, outlets and isolators, fabrication and/or modification of work bench and any other deemed necessary to ensure successful and safe installation and operation of the system should be included.
9.5	Must be able to utilise mains power supply of 220 - 240 V AC, 50 - 60 Hz. Otherwise, tenderer to include the necessary arrangements to ensure appropriate power supply required by the equipment.
10	WARRANTY
10.1	A minimum of two (2) years warranty for manufacturer's defect on the hardware, software and all cost of repairs and/or replacements should be included.
10.2	After-sales services and technical support shall continue to be provided by the Vendor beyond the warranty period.
10.3	One (1) on-site preventive maintenance to be carried out just before or soon after the one-year warranty period. Scope of work to follow manufacturer's manual / recommendation specific for the equipment offered, which includes but is not limited to: • Supply, delivery and installation of preventive maintenance kits and/or consumables • Software update (to obtain prior authorization from user and BME)

	• Inspection • Cleaning • Alignment • Calibration • Any other related preventive maintenance works required
11	DELIVERY
11.1	Items offered MUST be delivered within _____ from date of approval. (Vendor to indicate the delivery period.)
12	PRICE VALIDITY
12.1	Price validity MUST NOT BE LESS THAN one (1) year.

2. END USER TRAINING				
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)			
2	Tenderer must prepare a training attendance or proof of training done to end user during commissioning			

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

SECTION 3 - PROCUMENT DETAILS
BRAND:
MODEL:
DELIVERY TIME:
WARRANTY PERIOD:
COUNTRY OF ORIGIN:
LAST COUNTRY SOLD TO:
YEAR INTRODUCED TO MARKET:
PRICE VALIDITY: [AT LEAST ONE (1) YEAR PRICE VALIDTY]

SECTION 4 - TECHNICAL SPECIFICATION	
BRAND:	MODEL:
COUNTRY OF ORIGIN:	UNIT PRICE (B\$):
WARRANTY PERIOD:	TOTAL PRICE (B\$):
YEAR INTRODUCED TO MARKET:	LAST COUNTRY SOLD TO:
PRICE VALIDITY: [AT LEAST ONE (1) YEAR PRICE VALIDTY]	DELIVERY TIME:
AUTHORIZED DISTRIBUTOR: (AUTHORIZED DISTRIBUTOR LETTER ATTACHED)	
DETAILED BROCHURE INCLUDED	
USER AND SERVICE MANUALS:	
MAINS POWER SUPPLY:	
BATTERY	
POWER ADAPTER/CHARGER OUTPUT RATING:	
EQUIPMENT AMBIENT OPERATING TEMPERATURE RANGE:	
NUMBER OF TECHNICAL SUPPORT (ENGINEER/TECHNICIAN) Please provide training or certification for locals who is trained/certified	
DIMENSIONS AND WEIGHT OF MAIN UNIT:	
EQUIPMENT WHOLE LIFE TIME SUPPORT:	

SECTION 5 – 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 laboratory 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 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.
- _____ **time Planned Preventive Maintenance (PPM) PER YEAR** according to Manufacturer's Preventive Maintenance Guideline.

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.

	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 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	It is <u>MANDATORY</u> for the tenderer to do <u>site visit prior to tender submission</u> . Non-attendance will be considered as non-compliance .
6	Brochures / catalogues should be submitted / attached with tender document.
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)
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.

SECTION 3

TENDER FORM

TENDER REFERENCE NO: KK/243/2026/HTD

INVITATION TO TENDER

SUPPLY, DELIVERY, INSTALLATION, TESTING AND COMMISSIONING ONE (1) UNIT OF GENETIC ANALYSER SYSTEM FOR HUMAN IDENTIFICATION (HID) FOR FORENSIC BIOLOGY SECTION, DEPARTMENT OF SCIENTIFIC SERVICES, MINISTRY OF HEALTH

SECTION 1 – USER REQUIREMENTS				
1. STANDARD FEATURES				
NO.	DESCRIPTION	Tick (✓)		STATE OR SPECIFY OR REMARKS OR BROCHURE PAGE
		Y	N	
1.0	GENERAL			
1.1	Genetic Analyser System for Human Identification (HID) used to amplify forensic samples by polymerase chain reaction and perform fragment analysis by capillary electrophoresis (CE).			
1.2	This system shall comprise of the following: 1 unit of CE instrument. 2 units of thermal cycler. - Data management and acquisition system. - Accessories. - Performance check and training. - Consumables, reagents and kits for performance check and training. - Documentation of full developmental validation for forensic human identification use.			
2	CAPILLARY ELECTROPHORESIS INSTRUMENT SPECIFICATIONS			
2.1	Fluorescence-based capillary electrophoresis system.			
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2.14	Long-life, single-line 505nm, solid-state laser excitation source.			
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3	THERMAL CYCLER SPECIFICATIONS			
3.1	Block format: 96-well, 0.2ml alloy block			
3.2	Block range: 6 independent temperature zones (up to 10°C per zone), 30°C range across block			
3.3	Maximum block ramp rate: 6.0°C/sec			
3.4	Maximum sample ramp rate: 4.4°C/sec			
3.5	Temperature accuracy: $\pm 0.25^{\circ}\text{C}$ (35 – 99°C)			
3.6	Temperature range: 0 – 100.0°C			
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5	ACCESSORIES			
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5.2	Instrument to be supplied with all necessary standard operating accessories required to ensure successful operation.			
5.3	Two (2) units of low-bench working chair.			
6	METHOD VALIDATION & TRAINING			
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6.2	The performance check may include but is not limited to: • Standard Curve and Quality Metrics • Contamination Study • Minimum Threshold and Contamination Study • Sensitivity and Stochastic Study • Precision Study: Repeatability and Reproducibility • Accuracy Study: NIST Samples • Known and Non-Probative Sample Study • Mixture Study • Assessment of Non-Allelic Peaks			
6.3	On-site training for ALL staff members expected to handle the machine. Please ensure that adequate time is allocated such that training will take place in small groups to minimize staff shortage in the laboratory.			
6.4	Certificate of competence is to be issued to all trainees after completion of training.			
6.5	The successful tenderer needs to ensure • the key users are updated on the current and/or relevant information related to the system used. • provide ONE (1) off-site professional development in HID DNA analysis for TWO (2) key users. All expenses for attending the training shall be borne by the vendor; full registration, insurance, air ticket, daily allowance, accommodation and transport to and from the airport and training venue.			
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7.1	The successful tenderer needs to supply all necessary consumables, reagents and kits required for installation and commissioning, performance check and training.			
8	DOCUMENTATION & CERTIFICATION			
8.1	Operation manual (1 original and 1 copy).			
8.2	Availability of Developmental Validation paper documenting validation studies tested across Applied Biosystems™ STR PCR amplification kits and Precision ID NGS panels. based on requirements listed in the FBI Quality Assurance Standards for Forensic DNA Testing Laboratories and guidelines outlined by the Scientific Working Group on DNA Analysis Methods			
8.3	To be supplied during delivery/installation.			
9	SITE PREPARATION & ELECTRICAL REQUIREMENTS			
9.1	The tenderer shall ensure that the site preparation for the placement of the system taking into the consideration on the safety of the end user during operation of the instrument.			
9.2	It is MANDATORY for the tenderer to do site visit prior to tender submission to discuss site requirements.			

	A site visit form to be verified by user during the visit as evidence. Non-attendance will be considered as non-compliance .			
9.3	The site preparation details should be listed in the quotation / document submitted.			
9.4	All works involving additional electrical supply including wiring, outlets and isolators, fabrication and/or modification of work bench and any other deemed necessary to ensure successful and safe installation and operation of the system should be included.			
9.5	Must be able to utilise mains power supply of 220 - 240 V AC, 50 - 60 Hz. Otherwise, tenderer to include the necessary arrangements to ensure appropriate power supply required by the equipment.			
10	WARRANTY			
10.1	A minimum of two (2) years warranty for manufacturer's defect on the hardware, software and all cost of repairs and/or replacements should be included.			
10.2	After-sales services and technical support shall continue to be provided by the Vendor beyond the warranty period.			
10.3	One (1) on-site preventive maintenance to be carried out just before or soon after the one-year warranty period. Scope of work to follow manufacturer's manual / recommendation specific for the equipment offered, which includes but is not limited to: • Supply, delivery and installation of preventive maintenance kits and/or consumables • Software update (to obtain prior authorization from user and BME) • Inspection • Cleaning • Alignment • Calibration • Any other related preventive maintenance works required			
11	DELIVERY			
11.1	Items offered MUST be delivered within _____ from date of approval. (Vendor to indicate the delivery period.)			
12	PRICE VALIDITY			
12.1	Price validity MUST NOT BE LESS THAN one (1) year .			

2. END USER TRAINING				
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)			
2	Tenderer must prepare a training attendance or proof of training done to end user during commissioning			

SECTION 2 – PRICE PROPOSAL**UNIT PRICE:**
BND\$**TOTAL PRICE:**
BND\$**SECTION 3 - PROCUMENT DETAILS****BRAND:****MODEL:****DELIVERY TIME:****WARRANTY PERIOD:****COUNTRY OF ORIGIN:****LAST COUNTRY SOLD TO:****YEAR INTRODUCED TO MARKET:****PRICE VALIDITY:**[AT LEAST ONE (1) YEAR PRICE
VALIDITY]

SECTION 4 - 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:		220V-240V		OTHERS:	
		50-60HZ		OTHERS:	
BATTERY		RECHARGEABLE		☐	SINGLE-USE
		OTHERS:		☐	REPLACEABLE
	TYPE OF BATTERY:				
	RATING:				
POWER ADAPTER/CHARGER OUTPUT RATING:					
EQUIPMENT AMBIENT OPERATING TEMPERATURE RANGE:					
NUMBER OF TECHNICAL SUPPORT (ENGINEER/TECHNICIAN)	LOCAL		☐	☐ Trained / Certified	
Please provide training or certification for locals who is trained/certified	OVERSEA (SPECIFY LOCATION)		☐	☐ Not yet trained on the product	
DIMENSIONS AND WEIGHT OF MAIN UNIT:		☐ mm		☐	Kilogram (Kg)
		☐ cm		☐	Gram(g)
		☐ inch		☐	Pound (lbs)
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)				

SECTION 5 – 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 laboratory 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 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.
- _____ **time Planned Preventive Maintenance (PPM) PER YEAR** according to Manufacturer's Preventive Maintenance Guideline.

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

	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	It is MANDATORY for the tenderer to do <u>site visit prior to tender submission</u> . Non-attendance will be considered as non-compliance .	☐ Yes ☐ No
6	Brochures / catalogues should be submitted / attached with tender document.	☐ Yes ☐ No
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)	
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.	

SITE VISIT FORM

TENDER DETAILS		
TENDER REFERENCE	:	
TENDER TITLE	:	SUPPLY, DELIVERY, INSTALLATION, TESTING AND COMMISSIONING ONE (1) UNIT OF GENETIC ANALYSER SYSTEM FOR HUMAN IDENTIFICATION (HID) FOR FORENSIC BIOLOGY SECTION, DEPARTMENT OF SCIENTIFIC SERVICES, MINISTRY OF HEALTH.

COMPANY DETAILS				
COMPANY NAME	:		COMPANY STAMP AND SIGNATURE	
FOCAL PERSON	:			
		(Name, Designation)		
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	:		VERIFIED BY:	(Name, Designation, Signature)
DATE OF SITE VISIT	:			

Note: The Contractor **must visit the site before quoting any price** for the work stated in this Tender and shall satisfy himself as to the nature of work and site condition. The Contractor must fill in this form and obtain signature from the **user** as verification for having visited the Site.

Failing to do so will lead to **disqualification** from this Tender.

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 no 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