#### Rujukan Kami: (86) MOH/HQ/P/IKLAN-SH/2025

#### LAMPIRAN 4

BIL	Quotation Reference	Description	Advertisement Date	Closing Date (Not Later Than 09.00AM)	Quotation Fee	Requesting Department
4	273/CPC/2025/IKLAN/LAB	SUPPLY, DELIVERY, INSTALLATION, TESTING, AND COMMISSIONING OF ROTARY MICROTOME FOR HISTOLOGY LAB, DEPARTMENT OF LABORATORY SERVICES MINISTRY OF HEALTH.	10/06/2025	05/07/2025	\$5.00	JABATAN TEKNOLOGI PENJAGAAN KESIHATAN, KEMENTERIAN KESIHATAN

	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 <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)</b> 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	<b>DELIVERY PERIOD:</b> (Please state) Not More Than <b>90 days</b> 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.	
11	The vendor is required to provide proof of manufacture date confirming the equipment is new.	
12	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			
Please ☑ Tick where appropriate	Yes	No	Remarks
Supply of <u>One unit</u> of Rotary Microtome			

	3FL 11139 1 - U3FR REDBUREDUE			या किर्मान्त्री स्थान प्रशासकार हो। सक्ती राजिया का अस्ति स्थान स्थान
lease E	1 Tick where appropriate	Yes	No	Remarks
1	Type of Microtome: Manual or Semi-Motorized rotary microtome			Please specify:
2	Lockable, counter-balanced hand wheel for smooth and controlled movement.			
3	Includes a coarse feed wheel for initial coarse adjustments during sectioning.			
4	Specimen Retraction: Ability to perform specimen retraction for easier blade setup and retraction			
5	Section Thickness range up to 60 μm or 100 μm [Please specify range]			
6	Specimen Stroke Length: Vertical stroke length/cutting stroke capacity within range 64 - 72 mm			
7	Comes with a universal cassette clamp capable of securely holding most standard histological tissue cassettes.			
8	Adjustable object head allowing fine X/Y orientation of the block to be sectioned, ensuring precise alignment.			
9	Includes an adjustable blade block/holder to accommodate <b>both</b> high-profile and low-profile disposable blades.			
10	Accessories			
10.1	Supplied with essential standard tools for it to be operational			
10.2	A removable waste tray cover to collect tissue waste and prevent contamination.			

10.3	Dust cover included to protect the microtome when not in use		
11	Vendor shall perform a performance verification upon commissioning and ensure the performance of the installed equipment is within the acceptable limit of performance or as per manufacturer's recommendations or as per User's acceptance criteria. Vendor shall submit a copy of user verified Performance Qualification Report.		

2	END-USER TRAINING			
Please E	Tick where appropriate	Yes	No	Remarks
2.1	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)  Certificate of attendance and competence shall be issued to all trainees after completion of training.			
2.2	Tenderer must prepare a training attendance or proof of training done to end user during commissioning and the refresher course (6) months after commissioning.			

3	TECHNICAL TRAINING			
Please 5	☑ Tick where appropriate	Yes	No	Remarks
3.1	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)			

4	WARRANTY			
Please E	7 Tick where appropriate	Yes	No	Remarks
4.1	Tenderer to include warranty period of at least two (2) years			
4.2	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  • Planned Preventive Maintenance during warranty (one of which includes battery replacement at the end of warranty period).			

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

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

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

SECTION 3 -	PROCUMENT A	ND TECHNIC	AL S	PECIFICATION							
AUTHORIZED DISTRIBUTOR:	APPOINTED	BRUNEI DISTR	IBUT	OR	n in anac						
(AUTHORIZED DISTRIBUTOR LETTER	PROCURE F	ROM	cor	MPANY NAME:							
ATTACHED)	OVERSEA A		COMPANY ORIGIN:								
	DISTRIBUTO										
DETAILED BROCHURE INCLUDED	YES	NO	☑ or specify where appropriate								
ali di Santa da Maria da Mari Maria da Maria da Ma					Tenderers to acknowledge that they must provide at least TWO sets of USER AND SERVICE						
USER AND SERVICE MANUALS:	YES	NO	manuals when applying commissioning form.								
			One Set for End User, One Set for BME. (Please								
			prov	vide hardcopy or sof	ftcopy)						
MAINS POWER SUPPLY:	220V-240V	OTHERS:									
WAINS FOWER SUFFLI:	50-60HZ	OTHERS:			:						
	RECHARGE	ABLE		SINGLE-USE	REPLACEABLE						
	OTHERS:										
BATTERY CONTROL OF THE CONTROL OF TH	TYPE OF BATTERY	•									
	RATING:										
POWER ADAPTER/CHARGER OUTPUT	RATING:										
EQUIPMENT AMBIENT OPERATING TE	MPERATURE RANG	<b>E:</b>									
		1 7 6	☑ Tick where appropriate								
			<ul><li>☐ US FDA Standard,</li><li>☐ European Union CE MARK,</li></ul>								
INTERNATIONAL SAFETY STANDARD											
Must comply to at least 1 safety Standa	ards and certification	n		☐ Australian TO ☐ Canadian CS	GA Standard, A Standard or						
(Please attached the copy of stated sta				☐ Japanese JIS							
			Others (Please specify):								
NUMBER OF TECHNICAL SUPPORT				Trains	ed / Certified						
(ENGINEER/TECHNICIAN)	LOCAL				et trained on the						
				produ							
Please provide training or	OVERSEA										
certification for locals who is	(SPECIFY LOCATIO	N)	NEA	DEST LOCATION.							
trained/certified				REST LOCATION:	☐ Kilogram						
DIMENSIONS AND		□ m	m n		(Kg)						
WEIGHT OF MAIN UNIT:			ch		☐ Gram(g)						
					☐ Pound (lbs)						
		•			ment are available						
EQUIPMENT WHOLE LIFE TIME	for a minimum of	-									
SUPPORT:	extending beyond	-	-	cle of the equipm	ient.						
	No of years:	(Please spe	CITY)								

#### 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.

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**COMPANY CHOP AND SIGNATURE**