

BIL.	Quotation Reference	Description	Advertisement Date	Closing Date (Not Later Than 2.00PM)	Quotation Fee	Requesting Department	Focal Person
1	PPM/PROC/2023/<50K/002(MIC)	<p>TO SUPPLY AND DELIVER BLOOD CULTURE BOTTLES AND CONSUMABLES FOR NATIONAL CLINICAL MICROBIOLOGY REFERENCE LABORATORY, DEPARTMENT OF LABORATORY SERVICES, MINISTRY OF HEALTH. (CLUSTERING)</p> <p><u>PLACE OF SUBMISSION:</u>            QUOTATION BOX (GROUND FLOOR)            MINISTRY OF HEALTH            COMMONWEALTH DRIVE            BANDAR SERI BEGAWAN, BB 3910            NEGARA BRUNEI DARUSSALAM</p>	07/02/2023	11/02/2023	\$5.00	JABATAN PERKHIDMATAN MAKMAL	DR MUHD HAZIQ FIKRY HJ MOMIN

**ITEM(S) SPECIFICATIONS FOR ADVERTISEMENT (ABOVE \$2000)**

QUOTATION/TENDER REFERENCE NO:	
QUOTATION/TENDER NAME	TO SUPPLY AND DELIVER <b>BLOOD CULTURE BOTTLES AND CONSUMABLES</b> FOR <b>NATIONAL CLINICAL MICROBIOLOGY REFERENCE LABORATORY</b> , DEPARTMENT OF LABORATORY SERVICES, MINISTRY OF HEALTH. (CLUSTERING)

(Delete whichever is not applicable)

USER'S REQUIREMENTS				VENDOR'S OFFER					
NO	ITEM DESCRIPTIONS AND SPECIFICATIONS	PACKAGING SIZE	TOTAL ESTIMATE USAGE / YEAR	ITEM DESCRIPTIONS AND SPECIFICATIONS	PART/ CATALOGUE NUMBER AND BRAND	PACKAGING SIZE	TOTAL QUANTITY OFFERED / YEAR	*COST PER UNIT (B\$)	TOTAL COSTS (B\$)
1	Aerobic blood culture bottle	100 pieces/box	31 boxes						
2	Anaerobic blood culture bottle	100 pieces/box	31 boxes						
3	Paediatric blood culture bottle	100 pieces/box	24 boxes						
4	Airway Needles / Subculture Units	100 pieces/box	10 boxes						

NO	SPECIFICATIONS AND REQUIREMENTS	VENDOR'S OFFER (PLEASE STATE)
1.0	<b>BLOOD CULTURE BOTTLES</b>	
1.1	Aerobic blood culture bottles are FDA approved for aerobic and facultative anaerobic microorganisms (bacteria and fungi) from blood and other sterile body fluids.	
1.2	Anaerobic blood culture bottles are FDA approved for the culture and recovery of anaerobic and facultative anaerobic microorganisms (bacteria and fungi) from blood and other sterile body fluids.	
1.3	Paediatric blood culture bottles are FDA approved for the culture and recovery of aerobic micro-organisms (mainly bacteria and yeast) from paediatric patient blood samples as low as 0.05ml.	
1.4	All three types of bottles must be colour-coded in a distinctively different colour from each other.	
1.5	The blood culture bottles use sensitive Colorimetric Detection technology and enables Delay Vial Entry (DVE) up to 48 hours.	
1.6	The blood culture bottles are to be able to be stored at room temperature (up to 30°C).	
1.7	The blood culture bottles are safe to autoclave, light weight to save cost of disposal and shatterproof, made out of multi-layer polycarbonate for enhanced safety and can be used in pneumatic transport system.	
1.8	The blood culture bottles <b>MUST</b> be compatible to be used in the existing BacT/Alert Virtuo® blood culture system used in National Clinical Microbiology Reference Laboratory, RIPAS Hospital.	
2.0	<b>STERILE AIRWAY NEEDLE/SUBCULTURE UNITS</b>	
2.1	Used for dispensing blood culture samples to facilitate subculturing blood culture bottles	
2.2	Single-use, disposable and are individually, sterile packaged.	
2.3	Must be compatible to be used with the blood culture bottles (Aerobic, Anaerobic and Paediatric) to facilitate subculturing bottles.	
3.0	<b>SERVICE AND AFTER SALES SUPPORT</b>	

3.1	All reagent test kits / consumables supplied throughout this tender shall have a minimum expiry date of <b>six (6) months on delivery</b> . Should the reagent or consumable be urgently needed, provision of a reagent test kit or consumable with expiry date of less than six (6) months should be first agreed by the User of the particular laboratory before delivery is made.	
3.2	Letter of Undertaking (LOU) shall be produced upon each delivery of test kit or consumable with expiry date of less than six (6) months and vendor shall declare in the LOU that unused, unopened, expired kits will be replaced accordingly. For items which are known to have short expiry date such as those containing red blood cells, list down all such items and vendor shall declare in this tender submission of such items and shall be exempted from submitting LOU upon delivery.	
3.3	<b>Staggered delivery every 2 weeks usage</b> directly to the User. Delivery of the reagent test kits should be according to user schedule. Supplier should have other alternative in the case where supplier cannot fulfil the delivery on time.	
3.4	Product inserts and MSDS shall be provided to the User.	
3.5	Certificate of Analysis (CoA) for every batch and delivery of the reagent and/or consumables shall be provided to the User	
3.6	Any defect and contaminations occurring along the line should be replaced by the next shipment or at the earliest shipment.	
3.7	User shall have the rights to refuse delivery of items that do not meet the acceptance criteria such as, but not limited to, the following: <ol style="list-style-type: none"> <li>1. Tampered or damaged box</li> <li>2. Leakage upon delivery</li> <li>3. Items stored pre-delivery not in accordance to manufacturer's instructions</li> <li>4. Expiry date not meeting requirement</li> </ol>	
3.8	User shall have the rights to return any items, and to be replaced at no extra cost, if found not meeting the acceptance criteria upon opening a pack such as, but not limited to, the following: <ol style="list-style-type: none"> <li>1. Tampered or damaged packaging</li> <li>2. Evident of leakage or damaged products</li> </ol>	

	<p>3. Expired products that are evidently less than the requirement mentioned in para 3.1 calculated from delivery date</p> <p>4. Leakage and/or contaminated upon delivery</p>	
3.9	Vendor shall submit samples of the offered items directly to the Users no later than 7 days after the Closing Date of this advertisement or as required by the Users.	
4.0	<b>FINANCIAL AGREEMENT</b>	
4.1	Supply of the test kit including reagents, consumables and/or accessories is based on the number of kits required in the Purchase Order according to an agreed schedule period as stated in para 3.3.	
4.2	Buffer stock of the test kit including reagents, consumables and accessories shall be available at the local representative as contingency.	
4.3	Should there be any discontinuity of reagents / consumables due to non-compliance in the manufacturing of reagents; the vendor must be able to provide an alternative so that the test requests / services are still available for the customers.	
4.4	<p><b>EXIT CLAUSE:</b> The tender contract shall be automatically terminated even though tender has not yet expired and this shall be in effect due to, but not limited to, the following:</p> <ol style="list-style-type: none"> <li>1. When the testing is no longer required or relevant i.e. test is obsolete, to the laboratory or the Department.</li> <li>2. When the item(s) set out in this tender is/are no longer required by the laboratory or the Department.</li> <li>3. When the approved budget allocation for this tender contract has been used up before the tender contract expires whereby a renewal of tender shall be submitted by the user for an open advertisement subject to approval by the Mini Tender Board (<i>Lembaga Tawaran Kecil</i>).</li> </ol>	
5	<p><b>DELIVERY PERIOD:</b> Preferably 4 – 8 weeks and no later than 12 weeks after issue of Purchase Order</p>	<p>(Yes / No) (If No, please specify)</p>
6	<b>PRICE VALIDITY:</b>	

<p>The quotation shall remain valid for 6 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).</p>	
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\* 6 months validity required for <\$50K or 12 months for >\$50K

(Delete whichever is not applicable)

<b>DELIVERY PERIOD AFTER PO ISSUED</b>	<b>4-8 weeks and no longer than 12 weeks</b>		
Lab/Section/Unit	NATIONAL CLINICAL MICROBIOLOGY REFERENCE LABORATORY		
Lab/Section/Unit Ref No.:	DLS/PU/MIC/2023/U50K/01_BLD.CS		
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PPM/PROC Ref No.	PPM/PROC/2023/<50K/002(MIC)		
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