

	<b>Republic of the Philippines</b> <b>Department of Health</b> <b>BAGUIO GENERAL HOSPITAL AND MEDICAL CENTER</b> <b>Baguio City</b>	
	<b>BIDS AND AWARDS COMMITTEE</b> <b>SUPPLEMENTAL / BID BULLETIN</b>	Form No.: MCC-BAC-013

Bid Bulletin No.: **2023-004-BAC III**

Date: February 10, 2023

This Bid Bulletin is issued to modify or amend the Bid Documents of:

**SUPPLY, DELIVERY, INSTALLATION AND COMMISSIONING OF CARDIAC DEDICATED  
ULTRASOUND MACHINE**

**IB 2023-005 BAC III**

ORIGINAL	AMENDED
<b>Section III. BID DATA SHEET</b>	
<p><b>20.2</b> For purposes of Post-Qualification, the following document(s) shall be required. Within a non-extendible period of five (5) calendar days from receipt by the bidder of the notice from the BAC that it submitted the Lowest Calculated Bid (LCB), the Bidder shall submit the following documentary requirements:</p> <ol style="list-style-type: none"> <li>1. Latest Income and Business Tax returns filed and paid through the BIR Electronic Filing and Payment System (eFPS)</li> <li>2. License to Operate (LTO) if applicable.</li> <li>3. Warranty Certificate, if applicable;</li> <li>4. FDA Certificates (CPR/CMDN and CMDR whichever is applicable)</li> <li>5. Brochure (original or internet download), Technical Data Sheet and/or equivalent document of the product being offered showing compliance with the technical specifications: (If not in English, please refer to Clause 11 of the Instruction to Bidders);</li> <li>6. To conduct Product Demonstration within 5 calendar days upon notice of Post-Qualification (if required).</li> </ol> <p><i>Failure to submit any of the post-qualification requirements on time, or finding against the veracity thereof, shall disqualify the bidder for award.</i></p>	<p><b>20.2</b></p> <ul style="list-style-type: none"> <li>• <b><u>For the Opening of Bids, bidders are required to submit a detailed breakdown of their offer. This may be placed under their Price Schedule or in a separate sheet(s) of paper.</u></b></li> <li>• For purposes of Post-Qualification, the following document(s) shall be required. Within a non-extendible period of five (5) calendar days from receipt by the bidder of the notice from the BAC that it submitted the Lowest Calculated Bid (LCB), the Bidder shall submit the following documentary requirements:</li> </ul> <ol style="list-style-type: none"> <li>1. Latest Income and Business Tax returns filed and paid through the BIR Electronic Filing and Payment System (eFPS)</li> <li>2. License to Operate (LTO) if applicable.</li> <li>3. Warranty Certificate, if applicable;</li> <li>4. FDA Certificates (CPR/CMDN and CMDR whichever is applicable)</li> <li>5. Brochure (original or internet download), Technical Data Sheet and/or equivalent document of the product being offered showing compliance with</li> </ol>

<p><i>Provided in the event that a finding against the veracity of any of the documents submitted is made, it shall cause the forfeiture of the bid security in accordance with Section 69 of the IRR of RA 9184.</i></p>	<p>the technical specifications: (If not in English, please refer to Clause 11 of the Instruction to Bidders); 6.To conduct Product Demonstration within 5 calendar days upon notice of Post-Qualification (if required).</p> <p><i>Failure to submit any of the post-qualification requirements on time, or finding against the veracity thereof, shall disqualify the bidder for award. Provided in the event that a finding against the veracity of any of the documents submitted is made, it shall cause the forfeiture of the bid security in accordance with Section 69 of the IRR of RA 9184.</i></p>
<b>Section VII. TECHNICAL SPECIFICATIONS</b>	
<b>TRANSDUCER TYPES:</b>	<b>TRANSDUCER TYPES:</b>
A. Adult echo probe 3D capable, installed and ready to use.	A. Adult echo probe 3D capable, installed and ready to use.
B. Neonatal echo probe, installed and ready to use.	B. Neonatal echo probe, installed and ready to use.
C. Linear probe for vascular studies, installed and ready to use.	C. Linear probe for vascular studies, installed and ready to use.
D. Adult TEE probe – 3D capable and ready to use	D. Adult TEE probe – 3D capable and ready to use
E. Curvilinear probe for renal and abdominal aorta <del>All probes must be compatible and interchangeable with the existing cardiac ultrasound machines</del>	E. Curvilinear probe for renal and abdominal aorta

**BIDS AND AWARDS COMMITTEE III**

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**Approved by:**

**SGD**  
**RICARDO B. RUNEZ, JR., MD, FPCS, MHA, CESE**  
Medical Center Chief II