Securities and Exchange Commission PHILIPPINES

BAGONG PILIPINAS

**BIDS AND AWARDS COMMITTEE**
**NON-IT GOODS AND SERVICES**

**BID BULLETIN NO. 1**

This Bid Bulletin is issued to amend/clarify items in the Bidding Documents for the **SUPPLY AND DELIVERY OF MEDICAL SUPPLIES, MEDICINES, AND EQUIPMENT FOR THE SEC HEADQUARTERS USERS, PUBLIC BIDDING NO. 2025-027.**

**A. AMENDMENTS**

| Reference | Requirements/Specifications | Amendment |
|---|---|---|
| Section I. Invitation to Bid<br><br>Page 12 | 7) Bids must be duly received by the Bids and Awards Committee (BAC) Secretariat through manual submission at the office address indicated below on or before **29 August 2025 (Friday), 10:00AM**. Late bids shall not be accepted. | 7) Bids must be duly received by the Bids and Awards Committee (BAC) Secretariat through manual submission at the office address indicated below on or before **29 August 2025 (Friday), ~~10:00AM~~ 11:00AM**. Late bids shall not be accepted. |
| Section I. Invitation to Bid<br><br>Page 12 | 9) Bid opening shall be on **29 August 2025 (Friday), 10:15AM** at MSD Conference Room, 7F The SEC Headquarters, 7907 Makati Ave., Salcedo Village, Bel-Air, Makati City and via Zoom application. Bids will be opened in the presence of the Bidders' representatives who choose to attend the activity. | 9) Bid opening shall be on **29 August 2025 (Friday), ~~10:15AM~~ 11:15AM** at MSD Conference Room, 7F The SEC Headquarters, 7907 Makati Ave., Salcedo Village, Bel-Air, Makati City and via Zoom application. Bids will be opened in the presence of the Bidders' representatives who choose to attend the activity. |
| Section III. Bid Data Sheet<br><br>Page 45 | BDS Clause 16.1: The Bid Security shall be in the form of a Bid Securing Declaration, and any of the following:<br><br>xxx | BDS Clause 16.1: The Bid Security shall be in the form of a Bid Securing Declaration, ~~and~~ **or** any of the following:<br><br>xxx |
| Section III. Bid Data Sheet<br><br>Page 47 | BDS Clause 27.1: Licenses and Permits Required:<br><br>Food and Drug Administration (FDA) and/or ISO Certification, whichever is applicable. | ~~Food and Drug Administration (FDA) and/or ISO Certification, whichever is applicable.~~ |

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|---|---|---|
| | | 1. Company's valid and current License to Operate from the Food and Drug Administration (FDA);<br><br>2. Medicine's Certificate of Product Registration (CPR) from FDA. (For Lot 1) |
| Section VII. Technical Specifications<br><br>Page 78-79<br><br>Lot 2: Supply and Delivery of Automated External Defibrillator | | Certificate of Authorized Distributorship from the manufacturer. |
| Section VII. Technical Specifications<br><br>Page 78-79<br><br>Lot 2: Supply and Delivery of Automated External Defibrillator | | Accessories: one (1) extra pair of pads and one (1) cabinet per unit. |
| Section VII. Technical Specifications<br><br>Page 78-79<br><br>Lot 2: Supply and Delivery of Automated External Defibrillator | | Training and Certification: The winning bidder must conduct a one (1) day free onsite AED training (training date must be coordinated with the end-user). A Certificate of Completion must be issued to all personnel who participate in the training. |

**B. CLARIFICATIONS**

| Requirements | Bidder's Clarification | Response |
|---|---|---|
| Lot 2 – Automated External Defibrillator<br><br>Authorized Distributor Certificate | We recommend requiring bidders to submit an Authorized Distributor Certificate from the manufacturer. This will help ensure reliable after-sales support, including extended warranty coverage, availability of genuine parts, and competent service. | Bidders shall submit a Certificate of Authorized Distributorship from the manufacturer. It must be included in the bid submission. |

| | | |
|---|---|---|
| Lot 2 – Automated External Defibrillator<br><br>Extra Pad and Cabinet Package | We suggest including, for each AED unit, one (1) additional set of pads to serve as a spare in case of emergency use. This can be stated in the technical requirements as:<br><br>"Inclusion of Accessories: one (1) extra pair of pads per unit, and one (1) AED cabinet with strobe lights and alarm per unit, including installation at designated locations." | The extra pair of pads and cabinet per unit shall be provided by the winning bidder. In addition, the installation at designated locations may be conducted by the winning bidder at no additional cost to SEC. |
| Lot 2 – Automated External Defibrillator<br><br>Section III. Bid Data Sheet<br><br>Licenses and Permits Required:<br><br>Food and Drug Administration (FDA) and/or ISO Certification, whichever is applicable. | We recommend requiring bidders to hold an updated and valid License to Operate (LTO) from the Food and Drug Administration (FDA) as a Medical Device Distributor/Wholesaler. This will ensure the bidder is legally authorized and qualified to supply medical devices such as AEDs. | The recommendation is granted. Please refer to Annex B for the revised Technical Specifications. |
| Lot 2 – Automated External Defibrillator<br><br>Training and Certification | We request the inclusion of FREE onsite AED training, to be conducted once for one (1) day only upon delivery (or on a scheduled date), to ensure SEC end-users are equipped with the knowledge to operate the AED correctly, even though it is designated for layman use. A Certificate of Completion should be issued to all personnel who participate in the training/orientation. | The request is granted. Please refer to Annex B for the revised Technical Specifications. |

Provisions in the bidding documents that are in conflict with this Bulletin are deemed revised/amended. For guidance and information of all concerned.

20 August 2025

**ARMANDO A. PAN, JR.**
Chairman, Bids and Awards Committee
for Non-IT Goods and Services
/mrs

**Lot No. 2: Supply and Delivery of Automated External Defibrillator**

| Item No. | Specification | | Statement of Compliance |
|---|---|---|---|
| 1 | **Minimum Requirements** | | **Brand and Model:** |
| | **Defibrillator** | | |
| | Operations | Fully automatic / Semi-Automatic | |
| | Waveform | Biphasic Truncated Exponential (BTE) | |
| | Range of selected Energy | 100 to 360 J (Adult)<br>10 to 100 J (child) | |
| | Energy default | 200-300-360 J (Adult)<br>50-70-100 J (Child) | |
| | Power on time | < 2 seconds | |
| | ECG analysis time | < 5 seconds | |
| | Charge time | 0 seconds (as device is pre- charge during ECG analysis) | |
| | Time from power on to shock ready | < 8 seconds | |
| | **User prompts** | | |
| | CPR coaching | Voice guide, CPR metronome | |
| | **Controls** | | |
| | Lid release ON/OFF | Control device power on/off | |
| | Adult/child mode switch | Switch to child mode for reduced energy and appropriate CPR guidance | |
| | **Electrode Pads** | | |
| | Type | Pre connected, disposable for adult/child | |
| | Shelf life | 5 years (from date of manufacture) | |
| | **Automatic Self-test** | | |
| | Auto test | Daily, weekly, monthly, quarterly | |
| | Status indicator | Visual indicators indicating system readiness | |
| | **Data storage** | | |
| | Events | Up to 500 events | |
| | Voice recording | Up to 1 hour | |
| | CPR data | Up to 5 hours | |
| | Self-test report | 1000 records | |
| | Data export | Through USB flash memory | |
| | **Communication** | | |
| | Language button | Optional features allow the user to switch between maximum 3 languages | |
| | **Battery** | | |
| | Type | Lithium manganese dioxide (Li/MnO2), disposable 4200 mAh | |

| | | | |
|---|---|---|---|
| | Standby life | 6 years (performing auto test every week, not in use, not sending self-test report)<br><br>5 years (performing auto test every day, not in use, not sending self-test report) | |
| | Capacity | Max 400 shocks at 200 J, max 200 shocks for 360 J | |
| | Replace battery | Minimum 10 shocks at 200 J and 30 minutes | |
| | **Physical characteristics** | | |
| | Dimension | 210 mm (w) x 286 mm (d) x 78 mm (h) | |
| | Weight | 2.0 kg ~ 2.3 kg | |
| | **Environmental** | | |
| | Dust/ water resistance | IP55 | |
| | Temperature | Operating: -5 to 50 °C<br>Short term storage: -30 to 70 °C for a maximum 7 days Long term storage: 15 to 35 °C | |
| | Humidity | Operating/storage: 5 to 95% (non - condensing) | |
| | Drop | 1.5 m | |
| | **Warranty** | | |
| | AED Unit | AED must be covered by warranty for a period of eight (8) years against factory defect, from the date of delivery. | |
| | Accessories | one (1) extra pair of pads and one (1) cabinet per unit. | |
| **2** | **Green Specifications:** Outer box/Carton box must be 100% recycled fiber, or must be recyclable in the Philippines. | | |
| **3** | Certificate of Authorized Distributorship from the manufacturer. | | |
| **4** | Training and Certification: The winning bidder must conduct a one (1) day free onsite AED training (training date must be coordinated with the end-user). A Certificate of Completion must be issued to all personnel who participate in the training. | | |

Note: **Bidders must state "Comply" or "Not Comply" for each specification and provide the corresponding performance parameters for offered equipment.** Each response must be supported by a clear documentary support in the bid and properly cross-referenced. Acceptable evidence includes unaltered manufacturer sales brochures, official specification sheets, product samples, independent test results, and similar documents.

Claims should be backed by documentary support. If the evidence contradicts the claim, the bid may be rejected. Any false statement—whether in the compliance form or supporting documents—found during evaluation, post-qualification, or contract implementation may be considered fraudulent in accordance with

ITB Clause 3.1(a)(ii) and without prejudice to the imposition of appropriate administrative, civil, and criminal penalty in accordance with law.

I hereby certify that the statement of compliance with the foregoing technical specifications is true and correct.

Authorized Representative:

Signature : _____
Printed Name : _____
Company Name : _____
Date : _____
Contact Number : _____
Email Address : _____