



# National Center for Mental Health

## SUPPLEMENTAL BID BULLETIN

### ADDENDUM NO. 1

### PUBLIC BIDDING FOR THE SUPPLY AND DELIVERY OF DRUGS AND MEDICINES (SERVICE PATIENTS) CY 2024

This Supplemental Bid Bulletin No. 1 dated **November 21, 2023**, is issued to clarify, modify or amend items in the Bid Documents. This shall form an integral part of the Bid Documents.

ISSUES	CLARIFICATION
<p><b>LIST OF ITEMS</b></p> <p>ITEM CODE (DM-049) Norepinephrine 1 mg/ml, 4 ml ampule</p> <p>ITEM CODE (DM-149) Sterile Water for Injection 50ml QTY: 2,000 UOM: Vial</p> <p>ITEM CODE (DM-163) Glucose and Amino Acids with Electrolytes and Vitamin B1</p> <p>ITEM CODE (DM-167) Aztreonam 1 gm for injection QTY: 1,300 UOM: Ampule</p>	<p><b>LIST OF ITEMS</b></p> <p>ITEM CODE (DM-049) Norepinephrine 1 mg/ml, 4 ml ampule <b>*Stick with the NCMH Specifications</b></p> <p>ITEM CODE (DM-149) Sterile Water for Injection 50ml QTY: 2,000 UOM: <b>Bottle</b></p> <p>ITEM CODE (DM-163) Glucose and Amino Acids with Electrolytes and Vitamin B1, <b>1,000 ml or 1 liter</b></p> <p>ITEM CODE (DM-167) Aztreonam 1 gm for injection QTY: 1,300 UOM: <b>Vial</b></p>
<p><b>Section III BID DATA SHEET</b></p> <p><b>ITB Clause</b> <b>21.1</b> Additional contract documents relevant to the Project that is required by the Procuring Entity:</p> <p>xxx f. Certificate of Current Good Manufacturing Practice from FDA issued to Manufacturer.</p>	<p><b>Section III BID DATA SHEET</b></p> <p><b>ITB Clause</b> <b>21.1</b> Additional contract documents relevant to the Project that is required by the Procuring Entity:</p> <p>xxx f. Certificate of Current Good Manufacturing Practice from FDA issued to Manufacturer.</p>

*"There is no Health without Mental Health"*

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<p>****“Can we submit a Permit to Register issued by the Food and Drug Administration (FDA) in lieu of the Certificate of GMP Compliance (CGMP)?”</p>	<p><b>***For Foreign Drug Manufacturers, a Permit to Register issued by the FDA will be allowed in the absence of the certificate of GMP compliance.</b></p> <p><b><u>In accordance with FDA Circular No. 2020-020:</u></b></p> <p>IV. Guidelines  XX  XX  3. A Permit to Register shall be issued to the drug importer for applications found satisfactorily complying with the GMP standards based on document review.  XXX</p>
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**Other Matters:**

- A. Eligibility requirements and technical proposal should be in one folder and financial proposal in a separate folder, with shoelace on top or ring bound instead of a fastener, table of contents, and index tabs in words, not numbers.
- B. Folder of Eligibility requirements and technical proposal should be placed in one envelope. And the folder of the Financial proposal should be in another envelope. Both envelopes shall then be placed in one mother envelope marked as **“Original Bid”**
- C. Documents should be arranged chronologically according to the checklist issued.
- D. Color code for folders and envelope:  
**BLUE** – Public Bidding for the Supply and Delivery of Drugs and Medicines (Service Patients) CY 2024
- E. All other provisions on the bidding documents which are not affected shall remain in effect.
- F. The deadline for **Submission and Opening of Bids** is scheduled on **November 29, 2023 (Wednesday), 9:00 AM**, at the **Pag-Asa Hall**, National Center for Mental Health Compound, Mandaluyong City.
- G. Any bid submitted after the deadline for submission shall be declared **“LATE”** and shall NOT be accepted.
- H. The BAC shall open the bids immediately after the deadline for submission and receipt of bids.

For the information and guidance of all concerned.

**ALDEN C. CUYOS, MD, FPPA, IFAPA, MMHoA**  
Chairperson, BAC for Goods

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