Republic of the Philippines
Department of Health

# METRO MANILA CENTER FOR HEALTH DEVELOPMENT

## SUPPLEMENTAL/ BID BULLETIN NO. 1

**IB 2023 – 022**
**PROCUREMENT OF LEVONORGESTREL + ETHINYLESTRADIOL ORAL TABLETS "1 CYCLE PER BLISTER PACK (SHORT OF AWARD)**

This Supplemental/Bid Bulletin No. 1 is being issued to revise provisions/specifications in the Bidding Documents for a forecited project:

| **Revision and clarification to provisions/specifications in the Bidding Documents:** | |
|---|---|
| **Shelf Life**<br><br>Must be fresh commercial stock with a total shelf life of Forty eight (48) months upon manufacture and (42) months from the date of delivery.<br><br>- Not granted by the end-user | |
| **FROM** | **TO** |
| Shelf Life<br>1. Must be fresh commercial stock with a total shelf life of Sixty (60) months upon manufacture and (55) months from the date of delivery. | Shelf Life<br>1. Must be fresh commercial stock with a total shelf life of Sixty (60) months upon manufacture and (55) months from the date of delivery.<br>- Retained |

Bidders are advised to use the following attached forms and submit together with all required documents for the submission of bids on 28th day of November 2022, 9:30 AM:

This Supplemental/Bid Bulletin No. 1 shall form an integral part of the Bidding Documents. All other provisions indicated in the bidding documents which are not affected by this Supplemental/Bid Bulletin No. 1 shall remain in effect.

For guidance and information of all concerned.

Issued this 18th day of November 2022 in MMCHD

Approved by:

**ALELI ANNIE GRACE P. SUDIACAL, MD, MPH**
Director III / BAC Chairperson

Republic of the Philippines
Department of Health
**METRO MANILA CENTER FOR HEALTH DEVELOPMENT**

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| Republic of the Philippines<br>Department of Health<br>Metro Manila Center for Health Development<br><br>TECHNICAL SPECIFICATIONS | | | |
| Item No. 1 | **PROCUREMENT OF LEVONORGESTREL + ETHINYLESTRADIOL ORAL TABLETS "1 CYCLE PER BLISTER PACK (SHORT OF AWARD)** | Qty./Unit | 304,038 CYCLE |
| Name of Manufacturer: | | Country of Origin | |
| Brand: | | Model: (if applicable) | |
| **ABC: P 6,384,798.00** | | | |
| PURCHASER'S SPECIFICATION | | STATEMENT OF COMPLIANCE | |
| **SPECIFICATIONS:**<br><br>>oral tablets coated<br>>30 microgram Ethinylestradiol + 150 microgram Levonorgestrel<br>>28 tablets per cycle<br><br>**Packaging Instructions:**<br>1 cycle per blister pack<br>**Secondary Packaging**<br>Standard packaging of the manufacturer as approved by PFDA including product insert or encryption/imprint inside the box<br>**Shelf Life**<br>1. Must be fresh commercial stock with a total shelf life of Sixty (60) months upon manufacture and (55) months from the date of delivery.<br><br>**Labelling Instructions:**<br>>Standard labeling instruction as approved by PFDA pursuant to Administrative Order No. 2016-0008<br>>On each box, the following shall be imprinted or stickered with non-removable or permanent sticker or label that is binding, and with residue and tearing, if removed: | | | |

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| >In addition to the labeling requirements of PFDA:<br><br>Philippine Government Property – Department of Health NOT FOR SALE<br>Date of Manufacture:<br>Date of Expiry:<br>Batch/Lot No.:<br><br>**Delivery Schedule:**<br>>60-90 calendar days (CD)<br>**Delivery Site:**<br>>DOH-MMCHD Pasig Warehouse<br><br>**Additional Technical Documents:**<br>a. Valid PFDA Certificate of Product Registration (CPR) or Valid Extension<br>b. PFDA License to Operate (LTO) for Drugs Distributors and Traders<br>c. Product insert/ Product Information<br>d. Certificate from the manufacturer/main distributor that Bidder is an authorized dealer/exclusive distributor of the product<br><br>**FDA Test Analysis:**<br>>The minimum number of the sample units required for each test analysis of delivered medicines shall be based on the PFDA Circular 2014-014 | |
| **ADDITIONAL REQUIREMENT TO BE SUBMITTED BY THE SINGLE/LOWEST CALCULATED BIDDER (SCB/LCB) AS PART OF POST QUALIFICATION:**<br><br>1. One (1) original sample of manufacturer's product to be submitted and returned after evaluation. The sample submitted and approved during the evaluation shall be the same item to be delivered upon award of contract. Prototype of the labelling instruction must be part of the sample submitted however, the technical specifications of the labelling instruction of the product must be complied upon delivery. | |