



Republic of the Philippines
Department of Health

METRO MANILA CENTER FOR HEALTH DEVELOPMENT

SUPPLEMENTAL/ BID BULLETIN NO. 1

IB 2022 – 068

PROCUREMENT OF 1,852,283 TAB OF PARACETAMOL 500 MG TABLET

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:
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*No changes as stipulated in technical specifications.
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Bidders are advised to use the following attached forms and submit together with all required documents for the submission of bids on 19th day of July 2022, 9:00 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 12th day of July 2022 in MMCHD

Approved by:

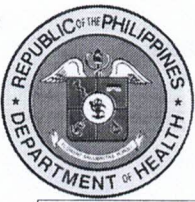
A handwritten signature in blue ink, appearing to read "Aleli", is written over the printed name.

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



Republic of the Philippines
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METRO MANILA CENTER FOR HEALTH DEVELOPMENT

Republic of the Philippines Department of Health Metro Manila Center for Health Development			
TECHNICAL SPECIFICATIONS			
Item No. 1	PROCUREMENT OF 1,852,283 TAB OF PARACETAMOL 500 MG TABLET	Qty./Unit	1,852,283 TABLETS
Name of Manufacturer:		Country of Origin	
Brand:		Model: (if applicable)	
ABC: P 1,000,232.82			
PURCHASER'S SPECIFICATION		STATEMENT OF COMPLIANCE	
<p>- 1,852,283 TAB OF PARACETAMOL 500 MG TABLET</p> <p><u>A. Upon delivery the following must be complied:</u></p> <p>-Shelf Life: Must be fresh commercial stock, with a total shelf life of 24 months from the date of manufacture but not less than 18 months from the date of delivery</p> <p>-Packaging Instructions: Standard packaging of the manufacturer approved by FDA</p> <p>-100 pieces per box</p> <p>-Labelling Instructions: Primary or Standard labelling instructions approved by FDA pursuant to AO No. 2016-0008</p> <p>In addition to the labelling requirements of the FDA:</p> <p>For the Secondary Labelling: each box should be imprinted or stickered with a permanent non-removable sticker/ label that is binding and will leave residue and ripping if removed</p> <p>Philippine Government Property-Department of Health NOT FOR SALE</p> <p>For the Tertiary Labelling: each box should be imprinted or stickered with a permanent non-removable sticker/ label that is binding and will leave residue and ripping if removed:</p> <p>Philippine Government Property-Department of Health NOT FOR SALE</p>			



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Date of Manufacture: _____

Date of Expiry: _____

Batch/Lot No. _____

Delivery Schedule: 30 to 45 Calendar days

Place of Delivery: Pasig Warehouse

D. Additional Requirement to be submitted by the Single/Lowest Calculated Bidder (SCB/LCB) as part of post qualification:

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.