

Republic of the Philippines DEPARTMENT OF HEALTH Metro Manila Center for Health Development



SUPPLEMENTAL/ BID BULLETIN NO. 2

2024-137 Procurement of Amlodipine 5 mg Tablet

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

Technical Specifications	
FROM	ТО
5mg as potassium salt, tablet	5mg as besilate/camsylate, tablet
5mg as potassium salt, tablet	5mg as besilate/camsylate, tablet

Bidders are advised to use the following attached forms and submit them together with all required documents for the submission of bids on the 22nd day of August 2024, 9:00 AM:

This Supplemental/Bid Bulletin No. 1 shall be integral to the Bidding Documents. All other provisions indicated in the bidding documents not affected by this Supplemental/Bid Bulletin No. 1 shall remain in effect.

For guidance and information of all concerned.

Issued this 19th day of August 2024 in MMCHD

Approved by:

JEREMIAS FRANCIS Y. CHAN, MD Licensing Officer V / BAC Chairperson

Republic of the Philippines Department of Health Metro Manila Center for Health Development

TECHNICAL SPECIFICATIONS

Item No. 1	Amlodipine 5 mg tablet	Qty./Unit	140,075 TPS	
Name of Manufacturer:		Country of Origin(if applicable)		
Brand:		Model: (if applicable)		
ABC: P 1	1,176,630.00			
PURCHASER'S SPECIFICATION		STATEMENT OF COMPLIANCE		
Specifica	tions:			
Route of	Administration: Oral			
Form and	Strength: 5mg as besilate/camsylate, tablet			
Delivery approved	Period: 30 – 60 calendar days from the receipt of NTP			

D. Upon delivery, the following shall be complied with:

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.

Packaging Instructions:

- 1. Primary Packaging: blister pack/foil strip
- 2. Secondary Packaging; 30 tablets per small box (DOH Treatment pack)
- 3. Tertiary Packaging: 1000 treatment packs per corrugated Carton

Labeling instructions:

Standard Labelling instruction as approved by FDA pursuant to Administrative Order no. 2016-008.

a. On each blister pack/foil strip, the following should be legibly imprinted or stickered:.

Philippine Government Property-Department of Health"

NOT FOR SALE

b. On each small/bigger box/corrugated carton, the foil owing should be imprinted or stickered with non removable or permanent sticker or label that is binding, with residue and tearing, if removed:

Philippine Government Property-Department of Health" NOT FOR SALE

Date of Manufacture:	
Date of Expiry:	
Batch/Lot No.	

B. Additional Requirements to be attached to Technical Specifications form arranged, numbered and tabbed as enumerated below:

1. Valid and current Certificate Product Registration (CPR) or Certificate of Medical Device

Registration (CMDR) or Certificate of Medical Device Notification (CMDN) issued by the Philippine Food and Drug Administration (PFDA)

The CPR must be valid for the entire period of the award. If the CPR/CMDN/CMDR is about to expire, the supplier must have submitted a copy of an application of renewal to the FDA at least 3 months before the expiry date (a copy of the expiring CPR, which is stamped with an "Extension of Validity" shall be submitted as proof); [AO 2019-0041]

- 2. Valid and current License to Operate (LTO) for Medical Device Exporter / Trader /Importer / Distributor / Wholesaler issued by Philippine Food and Drugs Administration (PFDA)
- 3.Product Insert/Product Information or downloaded from the internet and other manufacturer's unamended sales literature, unconditional statements of specification and compliance issued by the manufacturer, samples, independent test data, etc., as appropriate for a cross-referencing statement of compliance to the technical specification in accordance to what is indicated in Technical Specifications;
- 4. The bidder shall submit any of the following whichever is applicable:
- a. If the bidder is a manufacturer, certificate that the bidder manufactures the products/item or
- b. If the bidder is an Exclusive/Authorized Distributor or Dealer of the products/items, a Certificate or Contract from the manufacturer must be provided as proof that the bidder is an Exclusive/Authorized Distributor or Dealer of the products/items or
- c. If the bidder is an agent of the exclusive distributor or dealer, the following must be provided:
- i. Certificate or Distributor/Dealership Agreement by the Manufacturer with the distributor or dealer; and
- ii. Certificate/Contract between the distributor/dealer and the bidder
- 5. Valid Certificate of EPDMS