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

SUPPLEMENTAL/ BID BULLETIN NO. 1

IB 2022 - 049
PROCUREMENT OF LEVOTHYROXINE 50 MCG/25 MCG/ 12.5 MCG (LOT BIDDING)

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:
No Changes

Bidders are advised to use the following attached forms and submit together with all required documents for the submission of bids on 22nd day of March 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 15th day of March, 2022 in MMCHD

Approved by:

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

Section VI. Schedule of Requirements

The delivery schedule expressed as weeks/months stipulates hereafter a delivery date which is the date of delivery to the project site.

Item Number	Description	Quantity	Total ABC (Php)	Delivery Site	Delivered, Weeks/Months
	PROCUREMENT OF LEVOTHYROXINE 50 MCG TAB/ 25 MCG TAB/ 12.5 MCG TAB (LOT BIDDING)				
1	PROCUREMENT OF 85,000 PCS LEVOTHYROXINE 50 MCG TAB	85,000	510,000.00	DOH-MMCHD PASIG WAREHOUSE	30 CALENDAR DAYS
2	PROCUREMENT OF 100,000 PCS LEVOTHYROXINE 25 MCG TAB	100,000	500,000.00	DOH-MMCHD PASIG WAREHOUSE	30 CALENDAR DAYS
3	PROCUREMENT OF 125,000 PCS LEVOTHYROXINE 12.5 MCG TAB	125,000	500,000.00	DOH-MMCHD PASIG WAREHOUSE	30 CALENDAR DAYS

Technical Specifications

<table>
<tr><td colspan="4">Republic of the Philippines
Department of Health
Metro Manila Center for Health Development

TECHNICAL SPECIFICATIONS</td></tr>
<tr><td>Item No. 1</td><td>PROCUREMENT OF 85,000 PCS OF LEVOTHYROXINE 50 MCG TAB</td><td>Qty./Unit</td><td>85,000 pcs</td></tr>
<tr><td colspan="2">Name of Manufacturer:</td><td colspan="2">Country of Origin</td></tr>
<tr><td colspan="2">Brand:</td><td colspan="2">Model: (if applicable)</td></tr>
<tr><td colspan="2">ABC: 510,000.00</td><td colspan="2"></td></tr>
<tr><td colspan="2">PURCHASER'S SPECIFICATION</td><td colspan="2">STATEMENT OF COMPLIANCE</td></tr>
<tr><td colspan="2">A. Technical Specifications:

Form and Strength and Mode of Administration:
50 mcg tab/oral</td><td colspan="2"></td></tr>
<tr><td colspan="2">B. <u>Upon delivery the following must be complied:</u>
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
2. Standard packaging of the manufacturers as approved by the Philippine Food Drug Authority
Labelling Instructions:
Standard labelling instructions as approved by FDA pursuant to Administrative Order No. 2016-0008
In addition to the labelling requirements of the PFDA:
a. On each blister pack/foil strip and box, the following should be legibly imprinted or stickered using a permanent non-removable sticker/label that is binding and will leave residue and ripping if removed
b. On each small and bigger box/carton, the following should be legibly imprinted or stickered with non-removable or permanent sticker or label that is binding and will leave residue and ripping if removed

Philippine Government Property-Department of Health
NOT FOR SALE
Date of Manufacture: _____
Date of Expiry: _____
Batch/Lot No. _____
Delivery period: 30 Calendar days
Place of Delivery: DOH-MMCHD Pasig Warehouse</td><td colspan="2"></td></tr>
<tr><td colspan="2">C. <u>Additional Requirements to be attached with this form arranged, numbered and tabbed as enumerated below</u>:

1. Valid Certificate of Product Registration (CPR) issued by Philippine Food and Drug Administration (PFDA) or valid extension

The CPR must be valid for the entire period of the award. If the CPR is about to expire, the supplier must have submitted</td><td colspan="2"></td></tr>
</table>

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) as Medical Device Importer/ Wholesaler issued by PFDA. Provided that in case of expired LTO, the application for renewal was made timely as per PFDA Circular No. 2011-004. **In case of expired LTO, the following copies may be submitted:** a. Expired LTO; b. Application for renewal; and c. Official Receipt as proof of payment of renewal of LTO	
D. <u>Additional Requirement to be submitted by the Single/Lowest Calculated Bidder (SCB/LCB) as part of post qualification</u>: 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.	

Technical Specifications

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Metro Manila Center for Health Development

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<tr><td>Item No. 2</td><td>PROCUREMENT OF 100,000 PCS OF LEVOTHYROXINE 25 MCG TAB</td><td>Qty./Unit</td><td>100,000 pcs</td></tr>
<tr><td colspan="2">Name of Manufacturer:</td><td colspan="2">Country of Origin</td></tr>
<tr><td colspan="2">Brand:</td><td colspan="2">Model: (if applicable)</td></tr>
<tr><td colspan="2">ABC: 500,000.00</td><td colspan="2"></td></tr>
<tr><td colspan="2">PURCHASER'S SPECIFICATION</td><td colspan="2">STATEMENT OF COMPLIANCE</td></tr>
<tr><td colspan="2">A. Technical Specifications:

Form and Strength and Mode of Administration:
25 mcg tab/oral</td><td colspan="2"></td></tr>
<tr><td colspan="2">B. <u>Upon delivery the following must be complied:</u>
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
2. Standard packaging of the manufacturers as approved by the Philippine Food Drug Authority
Labelling Instructions:
Standard labelling instructions as approved by FDA pursuant to Administrative Order No. 2016-0008
In addition to the labelling requirements of the PFDA:
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Philippine Government Property-Department of Health
NOT FOR SALE
Date of Manufacture: _____
Date of Expiry: _____
Batch/Lot No. _____
Delivery period: 30 Calendar days
Place of Delivery: DOH-MMCHD Pasig Warehouse</td><td colspan="2"></td></tr>
<tr><td colspan="2">C. <u>Additional Requirements to be attached with this form arranged, numbered and tabbed as enumerated below</u>:

1 Valid Certificate of Product Registration (CPR) issued by Philippine Food and Drug Administration (PFDA) or valid extension

The CPR must be valid for the entire period of the award. If</td><td colspan="2"></td></tr>
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<tr><td>Item No. 3</td><td>PROCUREMENT OF 125,000 PCS OF LEVOTHYROXINE 12.5 MCG TAB</td><td>Qty./Unit</td><td>125,000 pcs</td></tr>
<tr><td colspan="2">Name of Manufacturer:</td><td colspan="2">Country of Origin</td></tr>
<tr><td colspan="2">Brand:</td><td colspan="2">Model: (if applicable)</td></tr>
<tr><td colspan="2">ABC: 500,000.00</td><td colspan="2"></td></tr>
<tr><td colspan="2">PURCHASER'S SPECIFICATION</td><td colspan="2">STATEMENT OF COMPLIANCE</td></tr>
<tr><td colspan="2">A. Technical Specifications:

Form and Strength and Mode of Administration:
12.5mcg tab/oral</td><td colspan="2"></td></tr>
<tr><td colspan="2">B. <u>Upon delivery the following must be complied:</u>
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
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Philippine Government Property-Department of Health
NOT FOR SALE
Date of Manufacture: _____
Date of Expiry: _____
Batch/Lot No. _____
Delivery period: 30 Calendar days
Place of Delivery: DOH-MMCHD Pasig Warehouse</td><td colspan="2"></td></tr>
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1 Valid Certificate of Product Registration (CPR) issued by Philippine Food and Drug Administration (PFDA) or valid extension

The CPR must be valid for the entire period of the award. If the CPR is about to expire, the supplier must have submitted a copy of an application of renewal to the FDA at least 3</td><td colspan="2"></td></tr>
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