

Republic of the Philippines Department of Budget and Management PROCUREMENT SERVICE BIDS AND AWARDS COMMITTEE



Bid Bulletin No. 1 November 20, 2018

Public Bidding 18-366-3

Supply and Delivery of Metformin for the Department of Health (DOH)

Issued pursuant to Sec. 22.5 of the IRR of R.A. 9184 to clarify, include and/or amend certain provisions in the Bidding Documents issued for this project, considering the issues raised and clarifications made by prospective bidders during the **Pre-Bid Conference** held on **13 November 2018**, likewise, respond to bidders' written queries received within the prescriptive period for filing.

A. AMENDMENT

BASES FOR CLARIFICATION/ REFERENCE AMENDMENT/INCLUSION SECTION I. INVITATION TO BID To clarify the item requirement page 4 for better understanding, see 1. XXX attached Appendix "B" Lot Section VII. Technical Item/Description Quantity Specifications No Treat METFORMIN, 500mg tablet or filmment 1,147,685 coated tablet Pack (TP)

2. The summary of the bidding activities is as follows:

Activities	Schedule
XXX	XXX
Last day of Submission of Written Clarification	November 15 <u>16</u> , 2018
Last day of Issuance of Bid Bulletin	November 19 <u>20</u> , 2018

To amend the schedule of bidding activities posted due to the postponement of the Prebid Conference on 13

November 2018, 8:30AM

For the purpose of this Bulletin and for better understanding of its contents, the following rules shall apply: (a) & Strikeout — denotes deletion; (b) <u>Underline</u> — denotes inclusion or new item/requirement; and "xxx" — denotes se phase/s being amended from the feet of the coain text. Street, Paco, Manila, Philippines 1007

Trunk line: (02) 689 7750 I helpdesk@ps-philgeps.gov.ph

Deadline of Submission of Bids	Tuesday, November 26 <u>27</u> , 2018, 1:30PM						
Opening of Bids	Immediately after the Deadline of Submission of Bids						

XXX

SECTION VI. SCHEDULE OF REQUIREMENTS

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Lot No.	Qt	y	Item/Description	Delivery period
No.	Qt	Treatm ent Pack (TP)	METFORMIN, 500mg tablet or film-coated tablet	Holding Area provided by the Supplier: Within Sixty (60) calendar days from the receipt date indicated in the Notice to Proceed (NTP), for the inspection and submission of samples for FDA testing; Delivery Period: Within 30 calendar days
				for the completion of deliveries to all access sites after the release of the
				"PASSED" result of FDA testing.

To amend the Schedule of Requirement, see attached Appendix "A"

Section VI. Schedule of Requirements.

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SUBMISSION OF SAMPLES

XXX

Lot No.	Item/Description	REQUIREMENTS
	METFORMIN, 500mg	Lay-out of the Packaging Design
1	tablet or film-coated tablet	(Attached as Appendix "C")

To clarify the item requirement for better understanding.

All deliveries shall be spursuant to GCC Claus shall include additional acceptance and testin	To amend the requirement for better understanding.	
	NICAL SPECIFICATIONS	
page 74 LOT NO.1 QUANTITY	: METFORMIN, 500mg tablet <u>or</u> : <u>film-coated tablet</u> : 1,147,685 Treatment Pack (TP)	To amend the form, see attached Appendix "B" Section VII.Technical Specifications
	BENCY SPECIFICATIONS ablet or film-coated tablet	

B. CLARIFICATIONS

REFERENCE	CLARIFICATION	RESOLUTION		
Section IIIBid-Data-Sheet ITB Clause 5.4 Page 39	Bidder clarified if the completed contract refers to paid contract.	Single Largest Completed Contract (SLCC) refers to paid and accepted contract.		
Section VI. Schedule of Requirements Page 71	See attached amendment for the Schedule of Requirements (refer to Appendix "A")			
Section VII. Technical Specifications.	Request to include the film-coated tablet.	See attached amendment for the Technical Specifications (refer to Appendix "B")		
Page 74-75	Request to consider the 168 tablets per Treatment Pack (TP)	No amendment.		

All other portions of the Bidding Documents affected by these amendments shall be made to conform to the same.

Amendments/inclusions/clarifications made herein shall be considered an integral part of the Bidding Documents.

The changes made in the Philippine Bidding Documents (5th Edition, August 2016) are deemed integrated in the terms and conditions for this project.

ENGR. EDWARD R. SADDI Chairperson, DBM-PS BAC 3

Section VI. Schedule of Requirements

lot				
No.	Qty		Item/Description	Delivery period
1	1,147,685	Treatment Pack (TP)	METFORMIN, 500mg tablet or film- coated tablet	Holding Area provided by the Supplier: Within Sixty (60) calendar days from the receipt date indicated in the Notice to Proceed (NTP), for the inspection and submission of samples for FDA testing; Delivery Period: Within 30 calendar days for the completion of deliveries to all access sites after the release of the "PASSED" result of FDA testing.

I hereby commit to deliver the required quality and quantities upon receipt of the Purchase Order as indicated above.

Name of Company	Signature Over Printed Name of	Date
	Authorized Representative	

N

SUBMISSION OF SAMPLES

All bidders shall submit the following samples/demo units to the Procurement Division 3 (PD 3) during post-qualification evaluation for evaluation and testing purposes:

Lot No.	Item/Description	REQUIREMENTS
		Lay-out of the Packaging Design
1	METFORMIN, 500mg tablet or film-coated tablet	(Attached as Appendix "C")

All deliveries shall be subject to Acceptance Test Requirements pursuant to GCC Clause 16 in which case the awardee-supplier shall include additional quantity in every batch of delivery for acceptance and testing purposes during FDA Test.

Evaluation of sample during post-qualification stage shall be conducted by the BAC, through joint PD 3 and DOH-TWG Representative using visual evaluation.

N

Section VII. Technical Specifications

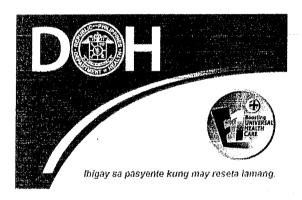
LOT NO 1		METFORMIN, 500mg tablet or film-coated tablet	
			Ī
QUANTITY	:	1,147,685 Treatment Pack (TP)	

AGENCY SPECIFICATIONS	BIDDER'S STATEMENT OF COMPLIANCE
METFORMIN, 500mg tablet or film-coated tablet	Brand:
Oral	
500mg tablet or film-coated tablet	
As Hydrochloride	
90 tablets per Treatment Pack	
REQUIRED DOCUMENTS:	
 Valid and Current Certificate of Product Registration Valid and current License to Operate (LTO) Certification from the manufacturer that the bidder is an Exclusive/Authorized Distributor or Dealer of the products/items, or a Distributorship/Dealership Agreement from the Manufacturer or Manufacturer's Agent or Main Distributor or Dealer in the Philippines. If the bidder is the manufacturer, certificate that the 	
bidder manufactures the products/items. Shelf Life:	
Must have a shelf life of at least twenty four (24) months from the date of manufacture but not less than eighteen (18) months from the date of delivery.	,
Packaging Instructions:	
Primary Packaging:	
Ninety (90) tablets per DOH Complete Treatment Pack	
Secondary Packaging: Standard packaging of the manufacturer including product package insert or encryption/imprint inside the box as approved by Philippine Food and Drug Administration (PFDA).	
Labeling Instructions:	
Standard labeling instructions as approved by PFDA pursuant to Administrative Order No. 2016-0008.	

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In addition to the labeling requirement of PFDA:	
On each blister pack/foil strip and treatment pack box, the following should be legibly imprinted or stickered with non-removable or permanent sticker/label that is binding and with residue and tearing, if removed:	
Philippine Government Property – DOH or Philippine Government Property –Department of Health	
NOT FOR SALE	
Manufacturing Date:	
Expiration Date:	
Batch/Lot No.:	
On each corrugated carton, the following should be legibly imprinted or stickered with non-removable or permanent sticker/label that is binding and with residue and tearing, if removed:	
 Philippine Government Property – DOH or Philippine Government Property –Department of Health	
NOT FOR SALE	
Manufacturing Date:	
Expiration Date:	
Batch/Lot No.:	
Product Recall/Replacement:	
 The supplier must ensure the quality of drugs and if there will be problems in the quality, the supplier will recall and replace the drugs distributed in the regions and health centers based on the Guidelines on Product recall, PFDA Circular No. 2016-012.	
We commit to deliver the goods under the new packaging a	as required by the Procurement Service.
I hereby certify that the statement of compliance to the for correct, otherwise, if found false either during evaluation of to automatic disqualification of our bid.	egoing technical specifications are true and post-qualification, the same shall give rise
Name of Company Signature Over Prin	ted Name Date
of Authorized Repre	esentative

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