



PHILIPPINE PHARMA PROCUREMENT, INC.  
Formerly PITC Pharma, Inc. (PPI)

**BIDS AND AWARDS COMMITTEE  
SUPPLEMENTAL/BID BULLETIN NO. 1**

**PROCUREMENT OF VARIOUS MEDICINE REQUIREMENTS FOR THE ARMED FORCES OF  
THE PHILIPPINES HEALTH SERVICE COMMAND (AFPHSC)  
UNDER ORDERING AGREEMENT  
(Project Ref. No. BAC-PO/GOODS 2018-08-001)**

This Supplemental Bid Bulletin is issued to clarify, modify and amend certain specifications/descriptions in the Invitation to Bid (ITB) and Bid Documents for the above-mentioned project.

I. The Schedule of Requirements (Section VI) is hereby amended to read as follows:

xxx                      xxx                      xxx  
G. Other Terms and Conditions:

xxx                      xxx                      xxx  
2. Terms and Conditions of Payment

xxx                      xxx                      xxx  
2.2. Required documents for payment are as follows:

2.2.1. BIR VAT registered Supplier's Invoice issued to PPPI

- **Supplier's Invoice to PPPI should be "PPPI for the account of AFPHSC"**

2.2.2. Supplier's Delivery Receipt duly received /signed by PPPI's authorized representative.

- **Supplier's Delivery Receipt duly received / signed by AFPHSC authorized representative, if applicable. However, only Delivery Receipt with Authority to Print (ATP) is acceptable.**

II. Additional Instruction in filling-up the Bidding Forms (Section IX):

A. The Letter of Intent (LOI) Form is revised to conform with the AFPHSC template. Please refer to the attached form. Please note that this form should be enclosed in the First Envelope prior to the TAB A requirement.

B. Additional instructions in filling-up Tab B of the Financial Component  
If there are columns that are not applicable in the computation of your cost, please indicate **"Not Applicable" or "N/A"**

III. Checklist of requirements (Section X) additional instruction are as follows:

A. Enclose the duly signed "Section VI. Schedule of Requirements" under the First Envelope, after Tab D.

B. Item No. 1.e under Post-Qualification Additional Requirements is deleted and no longer required

C. The following are included under Post-Qualification Additional Requirements and will now be as follows:

1. Valid and current License to Operate (LTO) as wholesaler, manufacturer or distributor with List of Sources issued by the Philippine Food and Drugs Administration (FDA) at the time of bidding and awarding and must be Certified True Copy by the Company Pharmacist.
2. Valid and current Certificate of Good Manufacturing Practice (cGMP) or in the case if foreign supplier / manufacturer, an equivalent document which must be authenticated by the Philippine Consulate and must be Certified True Copy by the Company Pharmacist.
3. Certification from the manufacturer that the bidder is an Authorized Distributor or Dealer of the products, or a Distributorship / Dealership Agreement from the manufacturer or manufacturer's Agent or Main Distributor or Dealer in the Philippines and must be Certified True Copy by the Company Pharmacist.

#### IV. Clarification on the requirement for Certificate of Analysis

The requirement on Certificate of Analysis (CoA) is different from the Results of Analysis (RA) issued by the Food and Drug Administration (FDA). The CoA required under this procurement should be issued by UST Center of Excellence in Drug Research Evaluation and Studies, Inc. (CEDRES) or UP-PGH for the following parenteral medicines:

- Cephalosporin First-Fourth Generation
- Sulbactam + Ampicillin Na
- Piperacillin +Tazobactam

**Please note that only the above mentioned medicines are required to submit Certificate of Analysis.**

**Note:** To avoid late submission of bids, bidders are encouraged to submit their bids as early as August 29, 2018.

This Supplemental Bid Bulletin shall form part of the ITB and the Bid Documents.

For the guidance and information of all concerned.

Issued this 23<sup>rd</sup> day of August 2018 in Makati City.



**JOSE A. CORTEZ**  
Chairperson  
Bids and Awards Committee