

PHILIPPINE BIDDING DOCUMENTS

(As Harmonized with Development Partners)

Procurement of GOODS

Government of the Republic of the Philippines

Name of the Project: “Pharmaceutical Supplies”

ITB No.: 072.21

*Procuring Entity: Philippine Heart Center
East Avenue, Quezon City*

Sixth Edition

July 2020

Preface

These Philippine Bidding Documents (PBDs) for the procurement of Goods through Competitive Bidding have been prepared by the Government of the Philippines for use by any branch, constitutional commission or office, agency, department, bureau, office, or instrumentality of the Government of the Philippines, National Government Agencies, including Government-Owned and/or Controlled Corporations, Government Financing Institutions, State Universities and Colleges, and Local Government Unit. The procedures and practices presented in this document have been developed through broad experience, and are for mandatory use in projects that are financed in whole or in part by the Government of the Philippines or any foreign government/foreign or international financing institution in accordance with the provisions of the 2016 revised Implementing Rules and Regulations of Republic Act No. 9184.

The Bidding Documents shall clearly and adequately define, among others: (i) the objectives, scope, and expected outputs and/or results of the proposed contract or Framework Agreement, as the case may be; (ii) the eligibility requirements of Bidders; (iii) the expected contract or Framework Agreement duration, the estimated quantity in the case of procurement of goods, delivery schedule and/or time frame; and (iv) the obligations, duties, and/or functions of the winning bidder.

Care should be taken to check the relevance of the provisions of the PBDs against the requirements of the specific Goods to be procured. If duplication of a subject is inevitable in other sections of the document prepared by the Procuring Entity, care must be exercised to avoid contradictions between clauses dealing with the same matter.

Moreover, each section is prepared with notes intended only as information for the Procuring Entity or the person drafting the Bidding Documents. They shall not be included in the final documents. The following general directions should be observed when using the documents:

- a. All the documents listed in the Table of Contents are normally required for the procurement of Goods. However, they should be adapted as necessary to the circumstances of the particular Procurement Project.
- b. Specific details, such as the “*name of the Procuring Entity*” and “*address for bid submission*,” should be furnished in the Instructions to Bidders, Bid Data Sheet, and Special Conditions of Contract. The final documents should contain neither blank spaces nor options.
- c. This Preface and the footnotes or notes in italics included in the Invitation to Bid, Bid Data Sheet, General Conditions of Contract, Special Conditions of Contract, Schedule of Requirements, and Specifications are not part of the text of the final document, although they contain instructions that the Procuring Entity should strictly follow.

- d. The cover should be modified as required to identify the Bidding Documents as to the Procurement Project, Project Identification Number, and Procuring Entity, in addition to the date of issue.
- e. Modifications for specific Procurement Project details should be provided in the Special Conditions of Contract as amendments to the Conditions of Contract. For easy completion, whenever reference has to be made to specific clauses in the Bid Data Sheet or Special Conditions of Contract, these terms shall be printed in bold typeface on Sections I (Instructions to Bidders) and III (General Conditions of Contract), respectively.

For guidelines on the use of Bidding Forms and the procurement of Foreign-Assisted Projects, these will be covered by a separate issuance of the Government Procurement Policy Board.

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Glossary of Acronyms, Terms, and Abbreviations

ABC – Approved Budget for the Contract.

BAC – Bids and Awards Committee.

Bid – A signed offer or proposal to undertake a contract submitted by a bidder in response to and in consonance with the requirements of the bidding documents. Also referred to as *Proposal* and *Tender*. (2016 revised IRR, Section 5[c])

Bidder – Refers to a contractor, manufacturer, supplier, distributor and/or consultant who submits a bid in response to the requirements of the Bidding Documents. (2016 revised IRR, Section 5[d])

Bidding Documents – The documents issued by the Procuring Entity as the bases for bids, furnishing all information necessary for a prospective bidder to prepare a bid for the Goods, Infrastructure Projects, and/or Consulting Services required by the Procuring Entity. (2016 revised IRR, Section 5[e])

BIR – Bureau of Internal Revenue.

BSP – Bangko Sentral ng Pilipinas.

Consulting Services – Refer to services for Infrastructure Projects and other types of projects or activities of the GOP requiring adequate external technical and professional expertise that are beyond the capability and/or capacity of the GOP to undertake such as, but not limited to: (i) advisory and review services; (ii) pre-investment or feasibility studies; (iii) design; (iv) construction supervision; (v) management and related services; and (vi) other technical services or special studies. (2016 revised IRR, Section 5[i])

CDA - Cooperative Development Authority.

Contract – Refers to the agreement entered into between the Procuring Entity and the Supplier or Manufacturer or Distributor or Service Provider for procurement of Goods and Services; Contractor for Procurement of Infrastructure Projects; or Consultant or Consulting Firm for Procurement of Consulting Services; as the case may be, as recorded in the Contract Form signed by the parties, including all attachments and appendices thereto and all documents incorporated by reference therein.

CIF – Cost Insurance and Freight.

CIP – Carriage and Insurance Paid.

CPI – Consumer Price Index.

DDP – Refers to the quoted price of the Goods, which means “delivered duty paid.”

DTI – Department of Trade and Industry.

EXW – Ex works.

FCA – “Free Carrier” shipping point.

FOB – “Free on Board” shipping point.

Foreign-funded Procurement or Foreign-Assisted Project– Refers to procurement whose funding source is from a foreign government, foreign or international financing institution as specified in the Treaty or International or Executive Agreement. (2016 revised IRR, Section 5[b]).

Framework Agreement – Refers to a written agreement between a procuring entity and a supplier or service provider that identifies the terms and conditions, under which specific purchases, otherwise known as “Call-Offs,” are made for the duration of the agreement. It is in the nature of an option contract between the procuring entity and the bidder(s) granting the procuring entity the option to either place an order for any of the goods or services identified in the Framework Agreement List or not buy at all, within a minimum period of one (1) year to a maximum period of three (3) years. (GPPB Resolution No. 27-2019)

GFI – Government Financial Institution.

GOCC – Government-owned and/or –controlled corporation.

Goods – Refer to all items, supplies, materials and general support services, except Consulting Services and Infrastructure Projects, which may be needed in the transaction of public businesses or in the pursuit of any government undertaking, project or activity, whether in the nature of equipment, furniture, stationery, materials for construction, or personal property of any kind, including non-personal or contractual services such as the repair and maintenance of equipment and furniture, as well as trucking, hauling, janitorial, security, and related or analogous services, as well as procurement of materials and supplies provided by the Procuring Entity for such services. The term “related” or “analogous services” shall include, but is not limited to, lease or purchase of office space, media advertisements, health maintenance services, and other services essential to the operation of the Procuring Entity. (2016 revised IRR, Section 5[r])

GOP – Government of the Philippines.

GPPB – Government Procurement Policy Board.

INCOTERMS – International Commercial Terms.

Infrastructure Projects – Include the construction, improvement, rehabilitation, demolition, repair, restoration or maintenance of roads and bridges, railways, airports, seaports, communication facilities, civil works components of information technology projects, irrigation, flood control and drainage, water supply, sanitation, sewerage and solid waste management systems, shore protection, energy/power and electrification facilities, national

buildings, school buildings, hospital buildings, and other related construction projects of the government. Also referred to as *civil works or works*. (2016 revised IRR, Section 5[u])

LGUs – Local Government Units.

NFCC – Net Financial Contracting Capacity.

NGA – National Government Agency.

PhilGEPS - Philippine Government Electronic Procurement System.

Procurement Project – refers to a specific or identified procurement covering goods, infrastructure project or consulting services. A Procurement Project shall be described, detailed, and scheduled in the Project Procurement Management Plan prepared by the agency which shall be consolidated in the procuring entity's Annual Procurement Plan. (GPPB Circular No. 06-2019 dated 17 July 2019)

PSA – Philippine Statistics Authority.

SEC – Securities and Exchange Commission.

SLCC – Single Largest Completed Contract.

Supplier – refers to a citizen, or any corporate body or commercial company duly organized and registered under the laws where it is established, habitually established in business and engaged in the manufacture or sale of the merchandise or performance of the general services covered by his bid. (Item 3.8 of GPPB Resolution No. 13-2019, dated 23 May 2019). Supplier as used in these Bidding Documents may likewise refer to a distributor, manufacturer, contractor, or consultant.

UN – United Nations.

Section I. Invitation to Bid

Notes on the Invitation to Bid

The Invitation to Bid (IB) provides information that enables potential Bidders to decide whether to participate in the procurement at hand. The IB shall be posted in accordance with Section 21.2 of the 2016 revised IRR of RA No. 9184.

Apart from the essential items listed in the Bidding Documents, the IB should also indicate the following:

- a. The date of availability of the Bidding Documents, which shall be from the time the IB is first advertised/posted until the deadline for the submission and receipt of bids;
- b. The place where the Bidding Documents may be acquired or the website where it may be downloaded;
- c. The deadline for the submission and receipt of bids; and
- d. Any important bid evaluation criteria (*e.g.*, the application of a margin of preference in bid evaluation).

The IB should be incorporated in the Bidding Documents. The information contained in the IB must conform to the Bidding Documents and in particular to the relevant information in the Bid Data Sheet.

Philippine Heart Center

INVITATION TO BID FOR “PHARMACEUTICAL SUPPLIES”

1. The *Philippine Heart Center (PHC)*, through the *PHC Corporate Operating Budget for CY 2022* intends to apply the sum of ***Php351,102,143.72*** being the ABC to payments under the contract for ***“Pharmaceutical Supplies”***. Bids received in excess of the ABC shall be automatically rejected at bid opening.

Select this for lot-procurement:

The *Philippine Heart Center (PHC)*, through the *PHC Corporate Operating Budget for CY 2022* intends to apply the sum of *Php351,102,143.72* being the ABC to payments under the contract for each lot/item. Bids received in excess of the ABC for each lot shall be automatically rejected at bid opening.

2. The *Philippine Heart Center (PHC)* now invites bids for the above Procurement Project. Delivery of the Goods is required by **60 Calendar Days**. Bidders should have completed, within ***the past two (2) years*** from the date of submission and receipt of bids, a contract similar to the Project. The description of an eligible bidder is contained in the Bidding Documents, particularly, in Section II (Instructions to Bidders).
3. Bidding will be conducted through open competitive bidding procedures using a non-discretionary *“pass/fail”* criterion as specified in the 2016 revised Implementing Rules and Regulations (IRR) of Republic Act (RA) No. 9184.

[Select one of the two following paragraphs and delete the other depending on the existence of conditions under Section 23.4.1.2 of the 2016 revised IRR of RA No. 9184]

- a. *[Select this paragraph if conditions (a), (c), and (d) under Section 23.4.1.2 of the 2016 revised IRR of RA No. 9184 do not exist:]* Bidding is restricted to Filipino citizens/sole proprietorships, partnerships, or organizations with at least sixty percent (60%) interest or outstanding capital stock belonging to citizens of the Philippines, and to citizens or organizations of a country the laws or regulations of which grant similar rights or privileges to Filipino citizens, pursuant to RA No. 5183.
4. Prospective Bidders may obtain further information from *Philippine Heart Center (PHC)* and inspect the Bidding Documents at the address given below during **8:00am to 5:00pm Monday to Friday**.
 5. A complete set of Bidding Documents may be acquired by interested Bidders on **October 15, 2021** from the given address and upon payment of the applicable fee for the Bidding Documents per Line Item. The Procuring Entity shall allow the bidder to

present its proof of payment for the fees in cash and *presented by the authorized person.*

[NOTE: For lot procurement, the maximum fee for the Bidding Documents for each lot shall be based on its ABC, in accordance with the Guidelines issued by the GPPB; provided that the total fees for the Bidding Documents of all lots shall not exceed the maximum fee prescribed in the Guidelines for the sum of the ABC of all lots.]

6. The *Philippine Heart Center (PHC)* will hold a Pre-Bid Conference¹ on *October 26, 2021, 9:00am* at *5th Floor, Amphitheater Conference Room, MAB Building, PHC*, which shall be open to prospective bidders.
7. Bids must be duly received by the BAC Secretariat through manual submission at the office address indicated below, on *November 09, 2021, 9:00am*. Late bids shall not be accepted.

PHC-Bids and Awards Committee
5th Floor, Amphitheater Conference Room
MAB Building, PHC

8. All Bids must be accompanied by a bid security in any of the acceptable forms and in the amount stated in **ITB** Clause 14.
9. Bid opening shall be on *November 09, 2021, 9:00am* at *5th Floor, Amphitheater Conference Room, MAB Building, PHC*. Bids will be opened in the presence of the bidders' representatives who choose to attend the activity.
10. *[Insert such other necessary information deemed relevant by the Procuring Entity such as the use of a back-up data or cloud storage for large files uploaded for online bid submissions]*
11. The *Philippine Heart Center (PHC)* reserves the right to reject any and all bids, declare a failure of bidding, or not award the contract at any time prior to contract award in accordance with Sections 35.6 and 41 of the 2016 revised IRR of RA No. 9184, without thereby incurring any liability to the affected bidder or bidders.
12. For further information, please refer to:

PHC-BAC Secretariat
Bids and Awards Committee
Philippine Heart Center
East Avenue, Quezon City
Telefax No. 925-24-01 local 4059
PHC Website: www.phc.gov.ph

¹ May be deleted in case the ABC is less than One Million Pesos (PhP1,000,000) where the Procuring Entity may not hold a Pre-Bid Conference.

13. You may visit the following websites:

For downloading of Bidding Documents: www.phc.gov.ph

MARIETTA A. VELASCO, RN, MAN
BAC Chairman for Pharmaceutical
Supplies and Medical Equipment

Section II. Instructions to Bidders

Notes on the Instructions to Bidders

This Section on the Instruction to Bidders (ITB) provides the information necessary for bidders to prepare responsive bids, in accordance with the requirements of the Procuring Entity. It also provides information on bid submission, eligibility check, opening and evaluation of bids, post-qualification, and on the award of contract.

1. Scope of Bid

The Procuring Entity, *Philippine Heart Center* wishes to receive Bids for the *Pharmaceutical Supplies*, with identification number *ITB.072.21*.

[Note: The Project Identification Number is assigned by the Procuring Entity based on its own coding scheme and is not the same as the PhilGEPS reference number, which is generated after the posting of the bid opportunity on the PhilGEPS website.]

The Procurement Project (referred to herein as “Project”) is composed of *[indicate number of lots or items]*, the details of which are described in Section VII (Technical Specifications).

2. Funding Information

2.1. The GOP through the source of funding as indicated below for *[indicate funding year]* in the amount of *[indicate amount]*.

2.2. The source of funding is:

- a. GOCC and GFIs, the proposed Corporate Operating Budget.

3. Bidding Requirements

The Bidding for the Project shall be governed by all the provisions of RA No. 9184 and its 2016 revised IRR, including its Generic Procurement Manuals and associated policies, rules and regulations as the primary source thereof, while the herein clauses shall serve as the secondary source thereof.

Any amendments made to the IRR and other GPPB issuances shall be applicable only to the ongoing posting, advertisement, or **IB** by the BAC through the issuance of a supplemental or bid bulletin.

The Bidder, by the act of submitting its Bid, shall be deemed to have verified and accepted the general requirements of this Project, including other factors that may affect the cost, duration and execution or implementation of the contract, project, or work and examine all instructions, forms, terms, and project requirements in the Bidding Documents.

4. Corrupt, Fraudulent, Collusive, and Coercive Practices

The Procuring Entity, as well as the Bidders and Suppliers, shall observe the highest standard of ethics during the procurement and execution of the contract. They or through an agent shall not engage in corrupt, fraudulent, collusive, coercive, and obstructive practices defined under Annex “I” of the 2016 revised IRR of RA No. 9184 or other integrity violations in competing for the Project.

5. Eligible Bidders

- 5.1. Only Bids of Bidders found to be legally, technically, and financially capable will be evaluated.
- 5.2. *[Select one, delete other/s]*
 - a. Foreign ownership exceeding those allowed under the rules may participate pursuant to:
 - i.i. When the Goods sought to be procured are not available from local suppliers; or
 - b. Foreign ownership limited to those allowed under the rules may participate in this Project.
- 5.3. Pursuant to Section 23.4.1.3 of the 2016 revised IRR of RA No.9184, the Bidder shall have an SLCC that is at least one (1) contract similar to the Project the value of which, adjusted to current prices using the PSA's CPI, must be at least equivalent to:
 - a. For the procurement of Expendable Supplies: The Bidder must have completed a single contract that is similar to this Project, equivalent to at least twenty-five percent (25%) of the ABC.
- 5.4. The Bidders shall comply with the eligibility criteria under Section 23.4.1 of the 2016 IRR of RA No. 9184.

6. Origin of Goods

There is no restriction on the origin of goods other than those prohibited by a decision of the UN Security Council taken under Chapter VII of the Charter of the UN, subject to Domestic Preference requirements under **ITB** Clause 18.

7. Subcontracts

- 7.a.1. The Bidder may subcontract portions of the Project to the extent allowed by the Procuring Entity as stated herein, but in no case more than twenty percent (20%) of the Project.

The Procuring Entity has prescribed that:

- 1.1.a. Subcontracting is not allowed.

8. Pre-Bid Conference

The Procuring Entity will hold a pre-bid conference for this Project on the specified date and time and either at its physical address *{[insert if applicable]}* and/or through videoconferencing/webcasting} as indicated in paragraph 6 of the **IB**.

9. Clarification and Amendment of Bidding Documents

Prospective bidders may request for clarification on and/or interpretation of any part of the Bidding Documents. Such requests must be in writing and received by the Procuring Entity, either at its given address or through electronic mail indicated in the **IB**, at least ten (10) calendar days before the deadline set for the submission and receipt of Bids.

10. Documents comprising the Bid: Eligibility and Technical Components

10.a.1. The first envelope shall contain the eligibility and technical documents of the Bid as specified in **Section VIII (Checklist of Technical and Financial Documents)**.

10.a.2. The Bidder's SLCC as indicated in **ITB** Clause 5.3 should have been completed within *the past 2 years* prior to the deadline for the submission and receipt of bids.

10.a.3. If the eligibility requirements or statements, the bids, and all other documents for submission to the BAC are in foreign language other than English, it must be accompanied by a translation in English, which shall be authenticated by the appropriate Philippine foreign service establishment, post, or the equivalent office having jurisdiction over the foreign bidder's affairs in the Philippines. Similar to the required authentication above, for Contracting Parties to the Apostille Convention, only the translated documents shall be authenticated through an apostille pursuant to GPPB Resolution No. 13-2019 dated 23 May 2019. The English translation shall govern, for purposes of interpretation of the bid.

11. Documents comprising the Bid: Financial Component

11.1. The second bid envelope shall contain the financial documents for the Bid as specified in **Section VIII (Checklist of Technical and Financial Documents)**.

11.2. If the Bidder claims preference as a Domestic Bidder or Domestic Entity, a certification issued by DTI shall be provided by the Bidder in accordance with Section 43.1.3 of the 2016 revised IRR of RA No. 9184.

11.3. Any bid exceeding the ABC indicated in paragraph 1 of the **IB** shall not be accepted.

11.4. For Foreign-funded Procurement, a ceiling may be applied to bid prices provided the conditions are met under Section 31.2 of the 2016 revised IRR of RA No. 9184.

12. Bid Prices

12.1. Prices indicated on the Price Schedule shall be entered separately in the following manner:

1.1.a. For Goods offered from within the Procuring Entity's country:

- i. The price of the Goods quoted EXW (ex-works, ex-factory, ex-warehouse, ex-showroom, or off-the-shelf, as applicable);
- ii. The cost of all customs duties and sales and other taxes already paid or payable;
- iii. The cost of transportation, insurance, and other costs incidental to delivery of the Goods to their final destination; and
- iv. The price of other (incidental) services, if any, listed in e.

1.1.b. For Goods offered from abroad:

- i. Unless otherwise stated in the **BDS**, the price of the Goods shall be quoted delivered duty paid (DDP) with the place of destination in the Philippines as specified in the **BDS**. In quoting the price, the Bidder shall be free to use transportation through carriers registered in any eligible country. Similarly, the Bidder may obtain insurance services from any eligible source country.
- ii. The price of other (incidental) services, if any, as listed in **Section VII (Technical Specifications)**.

13. Bid and Payment Currencies

13.a.1. For Goods that the Bidder will supply from outside the Philippines, the bid prices may be quoted in the local currency or tradeable currency accepted by the BSP at the discretion of the Bidder. However, for purposes of bid evaluation, Bids denominated in foreign currencies, shall be converted to Philippine currency based on the exchange rate as published in the BSP reference rate bulletin on the day of the bid opening.

13.a.2. Payment of the contract price shall be made in:

- 1.1.a. Philippine Pesos.

14. Bid Security

14.1. The Bidder shall submit a Bid Securing Declaration² or any form of Bid Security in the amount indicated in the **BDS**, which shall be not less than the percentage of the ABC in accordance with the schedule in the **BDS**.

14.2. The Bid and bid security shall be valid until *[indicate date]*. Any Bid not accompanied by an acceptable bid security shall be rejected by the Procuring Entity as non-responsive.

²In the case of Framework Agreement, the undertaking shall refer to entering into contract with the Procuring Entity and furnishing of the performance security or the performance securing declaration within ten (10) calendar days from receipt of Notice to Execute Framework Agreement.

15. Sealing and Marking of Bids

Each Bidder shall submit one copy of the first and second components of its Bid.

The Procuring Entity may request additional hard copies (**copy 1 and copy 2**) and/or electronic copies of the Bid. However, failure of the Bidders to comply with the said request shall not be a ground for disqualification.

ENVELOPE 1 shall contain the following Orange Folders in three separate envelopes properly marked as:

ORANGE FOLDER (1) – Original copies of Eligibility and Technical Documents as described in ITB Clause 20 (This should be enclosed in an envelope marked “*ORIGINAL ELIGIBILITY and TECHNICAL COMPONENT*”)

ORANGE FOLDER (2) – Original copies of Eligibility and Technical Documents as described in ITB Clause 20 (This should be enclosed in an envelope marked “*COPY NO.1 - ELIGIBILITY and TECHNICAL COMPONENT*”)

ORANGE FOLDER (3) – Original copies of Eligibility and Technical Documents as described in ITB Clause 20 (This should be enclosed in an envelope marked “*COPY NO.2 – ELIGIBILITY and TECHNICAL COMPONENT*”)

ENVELOPE 2 shall contain the following Yellow Folders in three separate envelopes properly marked as:

YELLOW FOLDER (1) – Original copies of the Financial Documents as described in ITB Clause 20 (This should be enclosed in an envelope marked “*ORIGINAL FINANCIAL COMPONENT*”)

YELLOW FOLDER (2) – Original copies of the Financial Documents as described in ITB Clause 20 (This should be enclosed in an envelope marked “*COPY NO. 1 - FINANCIAL COMPONENT*”)

YELLOW FOLDER (3) – Original copies of the Financial Documents as described in ITB Clause 20 (This should be enclosed in an envelope marked “*COPY NO. 2 - FINANCIAL COMPONENT*”)

16. Deadline for Submission of Bids

16.1. The Bidders shall submit on the specified date and time and either at its physical address or through online submission as indicated in paragraph 7 of the **IB**.

17. Opening and Preliminary Examination of Bids

17.1. The BAC shall open the Bids in public at the time, on the date, and at the place specified in paragraph 9 of the **IB**. The Bidders’ representatives who are present shall sign a register evidencing their attendance. In case videoconferencing, webcasting or other similar technologies will be used, attendance of participants shall likewise be recorded by the BAC Secretariat.

In case the Bids cannot be opened as scheduled due to justifiable reasons, the rescheduling requirements under Section 29 of the 2016 revised IRR of RA No. 9184 shall prevail.

- 17.2. The preliminary examination of bids shall be governed by Section 30 of the 2016 revised IRR of RA No. 9184.

18. Domestic Preference

- 18.1. The Procuring Entity will grant a margin of preference for the purpose of comparison of Bids in accordance with Section 43.1.2 of the 2016 revised IRR of RA No. 9184.

19. Detailed Evaluation and Comparison of Bids

- 19.1. The Procuring BAC shall immediately conduct a detailed evaluation of all Bids rated “*passed*,” using non-discretionary pass/fail criteria. The BAC shall consider the conditions in the evaluation of Bids under Section 32.2 of the 2016 revised IRR of RA No. 9184.
- 19.2. If the Project allows partial bids, bidders may submit a proposal on any of the lots or items, and evaluation will be undertaken on a per lot or item basis, as the case maybe. In this case, the Bid Security as required by **ITB** Clause 15 shall be submitted for each lot or item separately.
- 19.3. The descriptions of the lots or items shall be indicated in **Section VII (Technical Specifications)**, although the ABCs of these lots or items are indicated in the **BDS** for purposes of the NFCC computation pursuant to Section 23.4.2.6 of the 2016 revised IRR of RA No. 9184. The NFCC must be sufficient for the total of the ABCs for all the lots or items participated in by the prospective Bidder.
- 19.4. The Project shall be awarded as follows:

Option 2 – One Project having several items grouped into several lots, which shall be awarded as separate contracts per lot.

Option 3 - One Project having several items, which shall be awarded as separate contracts per item.
- 19.5. Except for bidders submitting a committed Line of Credit from a Universal or Commercial Bank in lieu of its NFCC computation, all Bids must include the NFCC computation pursuant to Section 23.4.1.4 of the 2016 revised IRR of RA No. 9184, which must be sufficient for the total of the ABCs for all the lots or items participated in by the prospective Bidder. For bidders submitting the committed Line of Credit, it must be at least equal to ten percent (10%) of the ABCs for all the lots or items participated in by the prospective Bidder.

20. Post-Qualification

- 20.2. Within a non-extendible period of five (5) calendar days from receipt by the Bidder of the notice from the BAC that it submitted the Lowest Calculated Bid, the Bidder shall submit its latest income and business tax returns filed and paid through the BIR Electronic Filing and Payment System (eFPS) and other appropriate licenses and permits required by law and stated in the **BDS**.

21. Signing of the Contract

- 21.1. The documents required in Section 37.2 of the 2016 revised IRR of RA No. 9184 shall form part of the Contract. Additional Contract documents are indicated in the **BDS**.

Section III. Bid Data Sheet

Notes on the Bid Data Sheet

The Bid Data Sheet (BDS) consists of provisions that supplement, amend, or specify in detail, information, or requirements included in the ITB found in Section II, which are specific to each procurement.

This Section is intended to assist the Procuring Entity in providing the specific information in relation to corresponding clauses in the ITB and has to be prepared for each specific procurement.

The Procuring Entity should specify in the BDS information and requirements specific to the circumstances of the Procuring Entity, the processing of the procurement, and the bid evaluation criteria that will apply to the Bids. In preparing the BDS, the following aspects should be checked:

- 18.a. Information that specifies and complements provisions of the ITB must be incorporated.
- 18.b. Amendments and/or supplements, if any, to provisions of the ITB as necessitated by the circumstances of the specific procurement, must also be incorporated.

Bid Data Sheet

ITB Clause	
5.3	For this purpose, contracts similar to the Project shall be: <ul style="list-style-type: none"> a. <i>Pharmaceutical Supplies.</i> b. completed within <i>the past 2 years</i> prior to the deadline for the submission and receipt of bids.
7.1	Not Applicable
12	The price of the Goods shall be quoted DDP [<i>state place of destination</i>] or the applicable International Commercial Terms (INCOTERMS) for this Project.
14.1	The bid security shall be in the form of a Bid Securing Declaration, or any of the following forms and amounts: <ul style="list-style-type: none"> a. The amount of not less than <i>Php7,022,042.87</i>, if bid security is in cash, cashier's/manager's check, bank draft/guarantee or irrevocable letter of credit; or b. The amount of not less than <i>Php17,555,107.19</i> if bid security is in Surety Bond.
19.3	<i>[In case the Project will be awarded by lot, list the grouping of lots by specifying the group title, items, and the quantity for every identified lot, and the corresponding ABC for each lot.]</i> <i>[In case the project will be awarded by item, list each item indicating its quantity and ABC.]</i>
20.2	<i>The Product should pass the scrutiny of the PHC-Therapeutic Committee</i>
21.2	<i>[List here any additional contract documents relevant to the Project that may be required by existing laws and/or the Procuring Entity.]</i>

Section IV. General Conditions of Contract

Notes on the General Conditions of Contract

The General Conditions of Contract (GCC) in this Section, read in conjunction with the Special Conditions of Contract in Section V and other documents listed therein, should be a complete document expressing all the rights and obligations of the parties.

Matters governing performance of the Supplier, payments under the contract, or matters affecting the risks, rights, and obligations of the parties under the contract are included in the GCC and Special Conditions of Contract.

Any complementary information, which may be needed, shall be introduced only through the Special Conditions of Contract.

1. Scope of Contract

This Contract shall include all such items, although not specifically mentioned, that can be reasonably inferred as being required for its completion as if such items were expressly mentioned herein. All the provisions of RA No. 9184 and its 2016 revised IRR, including the Generic Procurement Manual, and associated issuances, constitute the primary source for the terms and conditions of the Contract, and thus, applicable in contract implementation. Herein clauses shall serve as the secondary source for the terms and conditions of the Contract.

This is without prejudice to Sections 74.1 and 74.2 of the 2016 revised IRR of RA No. 9184 allowing the GPPB to amend the IRR, which shall be applied to all procurement activities, the advertisement, posting, or invitation of which were issued after the effectivity of the said amendment.

Additional requirements for the completion of this Contract shall be provided in the **Special Conditions of Contract (SCC)**.

2. Advance Payment and Terms of Payment

2.1. Advance payment of the contract amount is provided under Annex “D” of the revised 2016 IRR of RA No. 9184.

2.2. The Procuring Entity is allowed to determine the terms of payment on the partial or staggered delivery of the Goods procured, provided such partial payment shall correspond to the value of the goods delivered and accepted in accordance with prevailing accounting and auditing rules and regulations. The terms of payment are indicated in the **SCC**.

3. Performance Security

Within ten (10) calendar days from receipt of the Notice of Award by the Bidder from the Procuring Entity but in no case later than prior to the signing of the Contract by both parties, the successful Bidder shall furnish the performance security in any of the forms prescribed in Section 39 of the 2016 revised IRR of RA No. 9184.

4. Inspection and Tests

The Procuring Entity or its representative shall have the right to inspect and/or to test the Goods to confirm their conformity to the Project specifications at no extra cost to the Procuring Entity in accordance with the Generic Procurement Manual. In addition to tests in the **SCC, Section IV (Technical Specifications)** shall specify what inspections and/or tests the Procuring Entity requires, and where they are to be conducted. The Procuring Entity shall notify the Supplier in writing, in a timely manner, of the identity of any representatives retained for these purposes.

All reasonable facilities and assistance for the inspection and testing of Goods, including access to drawings and production data, shall be provided by the Supplier to the authorized inspectors at no charge to the Procuring Entity.

5. Warranty

- 6.1. In order to assure that manufacturing defects shall be corrected by the Supplier, a warranty shall be required from the Supplier as provided under Section 62.1 of the 2016 revised IRR of RA No. 9184.
- 6.2. The Procuring Entity shall promptly notify the Supplier in writing of any claims arising under this warranty. Upon receipt of such notice, the Supplier shall, repair or replace the defective Goods or parts thereof without cost to the Procuring Entity, pursuant to the Generic Procurement Manual.

6. Liability of the Supplier

The Supplier's liability under this Contract shall be as provided by the laws of the Republic of the Philippines.

If the Supplier is a joint venture, all partners to the joint venture shall be jointly and severally liable to the Procuring Entity.

Section V. Special Conditions of Contract

Notes on the Special Conditions of Contract

Similar to the BDS, the clauses in this Section are intended to assist the Procuring Entity in providing contract-specific information in relation to corresponding clauses in the GCC found in Section IV.

The Special Conditions of Contract (SCC) complement the GCC, specifying contractual requirements linked to the special circumstances of the Procuring Entity, the Procuring Entity's country, the sector, and the Goods purchased. In preparing this Section, the following aspects should be checked:

- a. Information that complements provisions of the GCC must be incorporated.
- b. Amendments and/or supplements to provisions of the GCC as necessitated by the circumstances of the specific purchase, must also be incorporated.

However, no special condition which defeats or negates the general intent and purpose of the provisions of the GCC should be incorporated herein.

Special Conditions of Contract

GCC Clause	
1	<p><i>[List here any additional requirements for the completion of this Contract. The following requirements and the corresponding provisions may be deleted, amended, or retained depending on its applicability to this Contract:]</i></p> <p>Delivery and Documents –</p> <p>For purposes of the Contract, “EXW,” “FOB,” “FCA,” “CIF,” “CIP,” “DDP” and other trade terms used to describe the obligations of the parties shall have the meanings assigned to them by the current edition of INCOTERMS published by the International Chamber of Commerce, Paris. The Delivery terms of this Contract shall be as follows:</p> <p><i>[For Goods supplied from abroad, state:]</i> “The delivery terms applicable to the Contract are DDP delivered <i>[indicate place of destination]</i>. In accordance with INCOTERMS.”</p> <p><i>[For Goods supplied from within the Philippines, state:]</i> “The delivery terms applicable to this Contract are delivered <i>[indicate place of destination]</i>. Risk and title will pass from the Supplier to the Procuring Entity upon receipt and final acceptance of the Goods at their final destination.”</p> <p>Delivery of the Goods shall be made by the Supplier in accordance with the terms specified in Section VI (Schedule of Requirements).</p> <p>For purposes of this Clause the Procuring Entity’s Representative at the Project Site is <i>[indicate name(s)]</i>.</p> <p>Incidental Services –</p> <p>The Supplier is required to provide all of the following services, including additional services, if any, specified in Section VI. Schedule of Requirements: <i>Select appropriate requirements and delete the rest.</i></p> <ul style="list-style-type: none"> a. performance or supervision of on-site assembly and/or start-up of the supplied Goods; b. furnishing of tools required for assembly and/or maintenance of the supplied Goods; c. furnishing of a detailed operations and maintenance manual for each appropriate unit of the supplied Goods; d. performance or supervision or maintenance and/or repair of the supplied Goods, for a period of time agreed by the parties, provided that this service shall not relieve the Supplier of any warranty obligations under this Contract; and

e. training of the Procuring Entity's personnel, at the Supplier's plant and/or on-site, in assembly, start-up, operation, maintenance, and/or repair of the supplied Goods.

f. [Specify additional incidental service requirements, as needed.]

The Contract price for the Goods shall include the prices charged by the Supplier for incidental services and shall not exceed the prevailing rates charged to other parties by the Supplier for similar services.

Spare Parts –

The Supplier is required to provide all of the following materials, notifications, and information pertaining to spare parts manufactured or distributed by the Supplier:

Select appropriate requirements and delete the rest.

2.i.1.a. such spare parts as the Procuring Entity may elect to purchase from the Supplier, provided that this election shall not relieve the Supplier of any warranty obligations under this Contract; and

2.i.1.b. in the event of termination of production of the spare parts:

i. advance notification to the Procuring Entity of the pending termination, in sufficient time to permit the Procuring Entity to procure needed requirements; and

ii. following such termination, furnishing at no cost to the Procuring Entity, the blueprints, drawings, and specifications of the spare parts, if requested.

The spare parts and other components required are listed in **Section VI (Schedule of Requirements)** and the cost thereof are included in the contract price.

The Supplier shall carry sufficient inventories to assure ex-stock supply of consumable spare parts or components for the Goods for a period of [*indicate here the time period specified. If not used indicate a time period of three times the warranty period*].

Spare parts or components shall be supplied as promptly as possible, but in any case, within [*insert appropriate time period*] months of placing the order.

	<p>Packaging –</p> <p>The Supplier shall provide such packaging of the Goods as is required to prevent their damage or deterioration during transit to their final destination, as indicated in this Contract. The packaging shall be sufficient to withstand, without limitation, rough handling during transit and exposure to extreme temperatures, salt and precipitation during transit, and open storage. Packaging case size and weights shall take into consideration, where appropriate, the remoteness of the Goods’ final destination and the absence of heavy handling facilities at all points in transit.</p> <p>The packaging, marking, and documentation within and outside the packages shall comply strictly with such special requirements as shall be expressly provided for in the Contract, including additional requirements, if any, specified below, and in any subsequent instructions ordered by the Procuring Entity.</p> <p>The outer packaging must be clearly marked on at least four (4) sides as follows:</p> <p>Name of the Procuring Entity Name of the Supplier Contract Description Final Destination Gross weight Any special lifting instructions Any special handling instructions Any relevant HAZCHEM classifications</p>
	<p>A packaging list identifying the contents and quantities of the package is to be placed on an accessible point of the outer packaging if practical. If not practical the packaging list is to be placed inside the outer packaging but outside the secondary packaging.</p> <p>Transportation –</p> <p>Where the Supplier is required under Contract to deliver the Goods CIF, CIP, or DDP, transport of the Goods to the port of destination or such other named place of destination in the Philippines, as shall be specified in this Contract, shall be arranged and paid for by the Supplier, and the cost thereof shall be included in the Contract Price.</p> <p>Where the Supplier is required under this Contract to transport the Goods to a specified place of destination within the Philippines, defined as the Project Site, transport to such place of destination in the Philippines, including insurance and storage, as shall be specified in this Contract, shall be arranged by the Supplier, and related costs shall be included in the contract price.</p>

	<p>Where the Supplier is required under Contract to deliver the Goods CIF, CIP or DDP, Goods are to be transported on carriers of Philippine registry. In the event that no carrier of Philippine registry is available, Goods may be shipped by a carrier which is not of Philippine registry provided that the Supplier obtains and presents to the Procuring Entity certification to this effect from the nearest Philippine consulate to the port of dispatch. In the event that carriers of Philippine registry are available but their schedule delays the Supplier in its performance of this Contract the period from when the Goods were first ready for shipment and the actual date of shipment the period of delay will be considered force majeure.</p> <p>The Procuring Entity accepts no liability for the damage of Goods during transit other than those prescribed by INCOTERMS for DDP deliveries. In the case of Goods supplied from within the Philippines or supplied by domestic Suppliers risk and title will not be deemed to have passed to the Procuring Entity until their receipt and final acceptance at the final destination.</p> <p>Intellectual Property Rights –</p> <p>The Supplier shall indemnify the Procuring Entity against all third-party claims of infringement of patent, trademark, or industrial design rights arising from use of the Goods or any part thereof.</p>
2.2	<p><i>[If partial payment is allowed, state]</i> “The terms of payment shall be as follows: _____.”</p>
4	<p>The inspections and tests that will be conducted are: <i>[Indicate the applicable inspections and tests]</i></p>

A. Determination of Lowest Calculated Bidder

B. Detailed Bid Evaluation (Eligibility and Technical Specification)

C. Post-Qualification

The post-qualification shall verify, validate, and ascertain all statements made and documents submitted by the bidder with the Lowest Calculated Bid/Highest Rated Bid, using non-discretionary criteria, as stated in the Bidding Documents. These criteria shall consider, but shall not be limited to, the following:

Class “A” Documents

Legal Documents

1. Valid PhilGEPS Registration Certificate (Platinum Membership) (all pages);
2. Registration certificate from Securities and Exchange Commission (SEC), Department of Trade and Industry (DTI) for sole proprietorship, or Cooperative Development Authority (CDA) for cooperatives or its equivalent document;
3. Mayor’s or Business permit issued by the city or municipality where the principal place of business of the prospective bidder is located, or the equivalent document for Exclusive Economic Zones or Areas;
4. Tax clearance per E.O. No. 398, s. 2005, as finally reviewed and approved by the Bureau of Internal Revenue (BIR).

Technical Documents

5. Statement of the prospective bidder of all its ongoing government and private contracts, including contracts awarded but not yet started, if any, whether similar or not similar in nature and complexity to the contract to be bid;
6. Statement of the bidder’s Single Largest Completed Contract (SLCC) similar to the contract to be bid, except under conditions provided for in Sections 23.4.1.3 and 23.4.2.4 of the 2016 revised IRR of RA No. 9184, within the relevant period as provided in the Bidding Documents;
7. Original copy of Bid Security. If in the form of a Surety Bond, submit also a certification issued by the Insurance Commission;
or
Original copy of Notarized Bid Securing Declaration;
8. Conformity with the Technical Specifications, must include production/ delivery schedule, manpower requirements, after-sales/parts.

9. Original duly signed Omnibus Sworn Statement (OSS);
and if applicable, Original Notarized Secretary's Certificate in case of a corporation, partnership, or cooperative; or Original Special Power of Attorney of all members of the joint venture giving full power and authority to its officer to sign the OSS and do acts to represent the Bidder.

Financial Documents

10. The Supplier's audited financial statements, showing, among others, the Supplier's total and current assets and liabilities, stamped "received" by the BIR or its duly accredited and authorized institutions, for the preceding calendar year which should not be earlier than two (2) years from the date of bid submission;
11. The prospective bidder's computation of Net Financial Contracting Capacity (NFCC);
or
A committed Line of Credit from a Universal or Commercial Bank in lieu of its NFCC computation.

Class "B" Documents

12. A duly signed joint venture agreement (JVA) in case the joint venture is already in existence;
or
duly notarized statements from all the potential joint venture partners stating that they will enter into and abide by the provisions of the JVA in the instance that the bid is successful.

Other documentary requirements under RA No. 9184 (as applicable)

13. *[For foreign bidders claiming by reason of their country's extension of reciprocal rights to Filipinos]* Certification from the relevant government office of their country stating that Filipinos are allowed to participate in government procurement activities for the same item or product.
14. Certification from the DTI if the Bidder claims preference as a Domestic Bidder or Domestic Entity.
15. Latest Income Tax Return (for monthly or quarterly tax remittance) *(Only tax returns filed and taxes paid through the BIR EFPS shall be accepted)*
16. Latest Business Tax (Percentage tax or VAT) Returns *(Only tax returns filed and taxes paid through the BIR EFPS shall be accepted)*
17. Certificate of Product Registration (CPR)
18. For those who submitted a Notarized Statement to form a Joint Venture during the Opening of Bids, the Joint Venture Agreement will be submitted with a disclosure of the percentage of ownership of both parties.

19. All parties to the Joint Venture shall submit the following:
 - a) Latest Income Tax Returns (for monthly or quarterly tax remittance) (*Only tax returns filed and taxes paid through the BIR EFPS shall be accepted*)
 - b) Latest Business Tax (Percentage tax or VAT) Returns (*Only tax returns filed and taxes paid through the BIR EFPS shall be accepted*)
 - c) Certificate of PhilGEPS Registration (Platinum Membership)

20. Certificate of Good Standing, Completion, **AND** Acceptance from PHC. Said Certification must be issued within the past twelve (12) months from bid submission. (This is applicable only to prospective bidder **with** previous contracts and completed projects with the PHC entered into within the past three (3) years from the submission and receipt of bids); **OR**

Duly Notarized Certificate of Good Standing, Completion, OR Acceptance from at least one (1) previous client. Said Certification must be issued within the past twelve (12) months from bid submission. (This is applicable only to prospective bidders **without** previous contracts and completed projects with the PHC.

21. A notarized certification to be duly submitted by the Supplier which provides that the Goods to be supplied under the contract are fresh/clean/pure/brand new stocks, appropriately sealed and labeled with an expiry date of not less than eighteen (18) months. In the case of biological drugs/products, the expiry date shall not be less than six (6) months from date of delivery.

Section VII. Technical Specifications

Notes for Preparing the Technical Specifications

A set of precise and clear specifications is a prerequisite for Bidders to respond realistically and competitively to the requirements of the Procuring Entity without qualifying their Bids. In the context of Competitive Bidding, the specifications (*e.g.* production/delivery schedule, manpower requirements, and after-sales service/parts, descriptions of the lots or items) must be prepared to permit the widest possible competition and, at the same time, present a clear statement of the required standards of workmanship, materials, and performance of the goods and services to be procured. Only if this is done will the objectives of transparency, equity, efficiency, fairness, and economy in procurement be realized, responsiveness of bids be ensured, and the subsequent task of bid evaluation and post-qualification facilitated. The specifications should require that all items, materials and accessories to be included or incorporated in the goods be new, unused, and of the most recent or current models, and that they include or incorporate all recent improvements in design and materials unless otherwise provided in the Contract.

Samples of specifications from previous similar procurements are useful in this respect. The use of metric units is encouraged. Depending on the complexity of the goods and the repetitiveness of the type of procurement, it may be advantageous to standardize the General Technical Specifications and incorporate them in a separate subsection. The General Technical Specifications should cover all classes of workmanship, materials, and equipment commonly involved in manufacturing similar goods. Deletions or addenda should then adapt the General Technical Specifications to the particular procurement.

Care must be taken in drafting specifications to ensure that they are not restrictive. In the specification of standards for equipment, materials, and workmanship, recognized Philippine and international standards should be used as much as possible. Where other particular standards are used, whether national standards or other standards, the specifications should state that equipment, materials, and workmanship that meet other authoritative standards, and which ensure at least a substantially equal quality than the standards mentioned, will also be acceptable. The following clause may be inserted in the Special Conditions of Contract or the Technical Specifications.

Sample Clause: Equivalency of Standards and Codes

Wherever reference is made in the Technical Specifications to specific standards and codes to be met by the goods and materials to be furnished or tested, the provisions of the latest edition or revision of the relevant standards and codes shall apply, unless otherwise expressly stated in the Contract. Where such standards and codes are national or relate to a particular country or region, other authoritative standards that ensure substantial equivalence to the standards and codes specified will be acceptable.

Reference to brand name and catalogue number should be avoided as far as possible; where unavoidable they should always be followed by the words “*or at least equivalent.*” References to brand names cannot be used when the funding source is the GOP.

Where appropriate, drawings, including site plans as required, may be furnished by the Procuring Entity with the Bidding Documents. Similarly, the Supplier may be requested to provide drawings or samples either with its Bid or for prior review by the Procuring Entity during contract execution.

Bidders are also required, as part of the technical specifications, to complete their statement of compliance demonstrating how the items comply with the specification.

Technical Specifications

[Bidders must state here either “Comply” or “Not Comply” against each of the individual parameters of each Specification stating the corresponding performance parameter of the equipment offered. Statements of “Comply” or “Not Comply” must be supported by evidence in a Bidders Bid and cross-referenced to that evidence. Evidence shall be in the form of manufacturer’s un-amended sales literature, unconditional statements of specification and compliance issued by the manufacturer, samples, independent test data etc., as appropriate. A statement that is not supported by evidence or is subsequently found to be contradicted by the evidence presented will render the Bid under evaluation liable for rejection. A statement either in the Bidder's statement of compliance or the supporting evidence that is found to be false either during Bid evaluation, post-qualification or the execution of the Contract may be regarded as fraudulent and render the Bidder or supplier liable for prosecution subject to the applicable laws and issuances.]

Directions in filling-up Schedule VII (Technical Specifications):

In filling-up the matrix on Statement of Compliance, the bidder shall provide relevant characteristics on each of the specific parameter such as its location in terms of the particular page, heading, and other provisions stated in the brochure, technical listing, operation manual, respectively.

To provide administrative ease in our evaluation, the bidder is required to provide a tab on each of the specific parameter (each correspondingly marked as Annex “A”, Annex “B”, etc.) for easy reference and validation purposes.

Item	Specifications Project: Pharmaceutical Supplies	Statement of Compliance
A	QUALIFICATIONS OF SUPPLIERS	
	1. A supplier whose drug/item(s) did not satisfactory pass all the evaluation stages of PHC-PTC, and hence, non-inclusive to the PHC's formulary, shall be ineligible to participate in the bidding process for the Project.	
	2. Only the most competent, responsible and duly accredited drug manufacturers/suppliers and/or exclusive distributors are qualified to participate in the public bidding. Exclusive Distributors shall submit, together with the bid proposals, an assurance from their principal/s that the quantities stipulated in the Purchase Order will be supplied on time. Participating Suppliers shall be registered with their quality control facilities duly accredited by the Food and Drug Administration (FDA).	
	3. Submission of Certificate of Product Registration (CPR).	
B	GENERAL CONDITIONS	
	1. The duration of the contract shall be within the prescribed period commencing on January 1, 2022 to December 31, 2022.	
	2. A supplier(s) is to bid on a per therapeutic category. The suppliers shall pay the corresponding amount of the bid documents according to their own preference to bid on a particular therapeutic category.	

Item	<p style="text-align: center;">Specifications Project: Pharmaceutical Supplies</p>	<p style="text-align: center;">Statement of Compliance</p>
	3. Antibiotics are subject to a Batch Certificate/Notification per Certificate of Product Registration (CPR).	
	4. The Supplier(s) shall ensure of fresh commercial stocks appropriately labeled. Expiry date shall not be less than eighteen (18) months and in the case of biological drugs/products shall be six (6) months from the date of delivery.	
	5. Bid prices shall be based on two decimal places only and shall be on a per unit/piece basis. The price quoted shall be final and fixed within the prescribed period of twelve (12) months without impositions of any new conditions and shall be inclusive of all applicable taxes. Pursuant to BIR Revenue Regulation No. 25-2018, "Providing for VAT Exemption on the Sale of Drugs and Medicines for Diabetes, High Cholesterol and Hypertension".	
	6. All price quotations per bid item shall be in typewritten form. Any corrections made on the price quotations shall likewise be typewritten and shall be properly initialed before the same is reproduced or photocopied (xerox). Non-compliance with this requirement shall cause automatic rejection of the bid offer.	
	7. If the supplier does not carry such bid item, indicate "NONE" in the corresponding space or cross the space. The exact packing or strength of bid item shall be clearly indicated in the price quotation.	
	8. A bid proposal which is qualified by the supplier through insertion or attachment of an unsolicited Terms of Conditions or Alternate Offer or any nature shall be rejected. Alternate offer means among other things, alterations of technical specifications or quantities called for a period of delivery required, or special offers (i.e., buy one take one).	
	9. The PHC reserves its right to procure bid items based on actual consumption due to medical exigencies that is determined by the PHC-PTC, and/or that demand advantageous to the interest of the PHC.	
C	DELIVERIES AND PENALTIES	
	1. Delivery of all items stated in the approved Purchase Order (P.O.) shall be strictly made within ten (10) calendar days from receipt.	
	2. Supplier of drugs/devices shall deliver only products registered under their names, except those with Certificate of Authorized/ Exclusive Distributorship.	
	3. A supplier shall warrant the delivery of high quality of the products that are free from all defects. The product/s to be delivered shall have a shelf life acceptable to the PHC and those which has expired shall be replaced and/or returned at no cost to the PHC or shall be automatically deducted one (1) month thereafter if without action from the supplier.	
	3.1 All deliveries in bottles and/or boxes shall be adequately sealed and labeled accordingly.	

Item	<p style="text-align: center;">Specifications Project: Pharmaceutical Supplies</p>	<p style="text-align: center;">Statement of Compliance</p>
	<p>3.2. Deliveries of antibiotics shall be accompanied by Batch Certificate and sensitivity discs (minimum of #3 cartridges per injectable antibiotic) for drug sensitivity testing by the PHC's Microbiology Laboratory.</p>	
	<p>3.3. All late deliveries/non-delivery in whole or in part shall be subject to the penalties prescribed under government accounting rules and regulation. The non-delivery of items shall result in the blacklisting of the supplier to participate in any future bidding in the PHC, and forfeiture of its Performance Security.</p>	
	<p>3.4. A supplier shall assure the PHC of a continuous supply of all items awarded. In cases where the supplier fails to deliver, the PHC shall procure from the second Lowest Calculated and Responsive Bidder. The price differential should be borne by the bid winner of the particular pharmaceutical item.</p>	

Section VIII. Checklist of Technical and Financial Documents

Notes on the Checklist of Technical and Financial Documents

The prescribed documents in the checklist are mandatory to be submitted in the Bid, but shall be subject to the following:

- a. GPPB Resolution No. 09-2020 on the efficient procurement measures during a State of Calamity or other similar issuances that shall allow the use of alternate documents in lieu of the mandated requirements; or
- b. Any subsequent GPPB issuances adjusting the documentary requirements after the effectivity of the adoption of the PBDs.

The BAC shall be checking the submitted documents of each Bidder against this checklist to ascertain if they are all present, using a non-discretionary “pass/fail” criterion pursuant to Section 30 of the 2016 revised IRR of RA No. 9184.

Checklist of Technical and Financial Documents

I. TECHNICAL COMPONENT ENVELOPE

Class “A” Documents

Legal Documents

- (1.1.a) Valid PhilGEPS Registration Certificate (Platinum Membership) (all pages);
- (1.1.b) Registration certificate from Securities and Exchange Commission (SEC), Department of Trade and Industry (DTI) for sole proprietorship, or Cooperative Development Authority (CDA) for cooperatives or its equivalent document;
- (1.1.c) Mayor’s or Business permit issued by the city or municipality where the principal place of business of the prospective bidder is located, or the equivalent document for Exclusive Economic Zones or Areas;
- (1.1.d) Tax clearance per E.O. No. 398, s. 2005, as finally reviewed and approved by the Bureau of Internal Revenue (BIR).

Technical Documents

- (f) Statement of the prospective bidder of all its ongoing government and private contracts, including contracts awarded but not yet started, if any, whether similar or not similar in nature and complexity to the contract to be bid;
- (g) Statement of the bidder’s Single Largest Completed Contract (SLCC) similar to the contract to be bid, except under conditions provided for in Sections 23.4.1.3 and 23.4.2.4 of the 2016 revised IRR of RA No. 9184, within the relevant period as provided in the Bidding Documents;
- (h) Original copy of Bid Security. If in the form of a Surety Bond, submit also a certification issued by the Insurance Commission;
or
Original copy of Notarized Bid Securing Declaration;
- (i) Conformity with the Technical Specifications, must include production/delivery schedule, manpower requirements, after-sales/parts.
- (j) Original duly signed Omnibus Sworn Statement (OSS);
and if applicable, Original Notarized Secretary’s Certificate in case of a corporation, partnership, or cooperative; or Original Special Power of Attorney of all members of the joint venture giving full power and authority to its officer to sign the OSS and do acts to represent the Bidder.

Financial Documents

- (k) The Supplier's audited financial statements, showing, among others, the Supplier's total and current assets and liabilities, stamped "received" by the BIR or its duly accredited and authorized institutions, for the preceding calendar year which should not be earlier than two (2) years from the date of bid submission;

- (l) The prospective bidder's computation of Net Financial Contracting Capacity (NFCC);
or
A committed Line of Credit from a Universal or Commercial Bank in lieu of its NFCC computation.

Class "B" Documents

- (m) A duly signed joint venture agreement (JVA) in case the joint venture is already in existence;
or
duly notarized statements from all the potential joint venture partners stating that they will enter into and abide by the provisions of the JVA in the instance that the bid is successful.

Or Certificate of Simplified Supplier's Registration (CSSR)

Other documentary requirements under RA No. 9184 (as applicable)

- (n) *[For foreign bidders claiming by reason of their country's extension of reciprocal rights to Filipinos]* Certification from the relevant government office of their country stating that Filipinos are allowed to participate in government procurement activities for the same item or product.

- (o) Certification from the DTI if the Bidder claims preference as a Domestic Bidder or Domestic Entity.

25 FINANCIAL COMPONENT ENVELOPE

- (a) Original of duly signed and accomplished Financial Bid Form;
- (b) Abstract of Sealed Bidding BID FORM
- (c) Original of duly signed and accomplished Price Schedule(s).

Omnibus Sworn Statement (Revised)

[shall be submitted with the Bid]

REPUBLIC OF THE PHILIPPINES)
CITY/MUNICIPALITY OF _____) S.S.

AFFIDAVIT

I, [Name of Affiant], of legal age, [Civil Status], [Nationality], and residing at [Address of Affiant], after having been duly sworn in accordance with law, do hereby depose and state that:

1. *[Select one, delete the other:]*

[If a sole proprietorship:] I am the sole proprietor or authorized representative of [Name of Bidder] with office address at [address of Bidder];

[If a partnership, corporation, cooperative, or joint venture:] I am the duly authorized and designated representative of [Name of Bidder] with office address at [address of Bidder];

2. *[Select one, delete the other:]*

[If a sole proprietorship:] As the owner and sole proprietor, or authorized representative of [Name of Bidder], I have full power and authority to do, execute and perform any and all acts necessary to participate, submit the bid, and to sign and execute the ensuing contract for [Name of the Project] of the [Name of the Procuring Entity], as shown in the attached duly notarized Special Power of Attorney;

[If a partnership, corporation, cooperative, or joint venture:] I am granted full power and authority to do, execute and perform any and all acts necessary to participate, submit the bid, and to sign and execute the ensuing contract for [Name of the Project] of the [Name of the Procuring Entity], as shown in the attached [state title of attached document showing proof of authorization (e.g., duly notarized Secretary's Certificate, Board/Partnership Resolution, or Special Power of Attorney, whichever is applicable)];

3. [Name of Bidder] is not "blacklisted" or barred from bidding by the Government of the Philippines or any of its agencies, offices, corporations, or Local Government Units, foreign government/foreign or international financing institution whose blacklisting rules have been recognized by the Government Procurement Policy Board, **by itself or by relation, membership, association, affiliation, or controlling interest with another blacklisted person or entity as defined and provided for in the Uniform Guidelines on Blacklisting;**

4. Each of the documents submitted in satisfaction of the bidding requirements is an authentic copy of the original, complete, and all statements and information provided therein are true and correct;

5. [Name of Bidder] is authorizing the Head of the Procuring Entity or its duly authorized representative(s) to verify all the documents submitted;

6. *[Select one, delete the rest:]*

[If a sole proprietorship:] The owner or sole proprietor is not related to the Head of the Procuring Entity, members of the Bids and Awards Committee (BAC), the Technical Working Group, and the BAC Secretariat, the head of the Project Management Office or the end-user unit, and the project consultants by consanguinity or affinity up to the third civil degree;

[If a partnership or cooperative:] None of the officers and members of *[Name of Bidder]* is related to the Head of the Procuring Entity, members of the Bids and Awards Committee (BAC), the Technical Working Group, and the BAC Secretariat, the head of the Project Management Office or the end-user unit, and the project consultants by consanguinity or affinity up to the third civil degree;

[If a corporation or joint venture:] None of the officers, directors, and controlling stockholders of *[Name of Bidder]* is related to the Head of the Procuring Entity, members of the Bids and Awards Committee (BAC), the Technical Working Group, and the BAC Secretariat, the head of the Project Management Office or the end-user unit, and the project consultants by consanguinity or affinity up to the third civil degree;

7. *[Name of Bidder]* complies with existing labor laws and standards; and

8. *[Name of Bidder]* is aware of and has undertaken the responsibilities as a Bidder in compliance with the Philippine Bidding Documents, which includes:

- a. Carefully examining all of the Bidding Documents;
- b. Acknowledging all conditions, local or otherwise, affecting the implementation of the Contract;
- c. Making an estimate of the facilities available and needed for the contract to be bid, if any; and
- d. Inquiring or securing Supplemental/Bid Bulletin(s) issued for the *[Name of the Project]*.

9. *[Name of Bidder]* did not give or pay directly or indirectly, any commission, amount, fee, or any form of consideration, pecuniary or otherwise, to any person or official, personnel or representative of the government in relation to any procurement project or activity.

10. In case advance payment was made or given, failure to perform or deliver any of the obligations and undertakings in the contract shall be sufficient grounds to constitute criminal liability for Swindling (Estafa) or the commission of fraud with unfaithfulness or abuse of confidence through misappropriating or converting any payment received by a person or entity under an obligation involving the duty to deliver certain goods or services, to the prejudice of the public and the government of the Philippines pursuant to Article 315 of Act No. 3815 s. 1930, as amended, or the Revised Penal Code.

IN WITNESS WHEREOF, I have hereunto set my hand this ___ day of ___, 20__ at _____, Philippines.

*[Insert NAME OF BIDDER OR ITS AUTHORIZED
REPRESENTATIVE]
[Insert signatory's legal capacity]
Affiant*

[Jurat]
[Format shall be based on the latest Rules on Notarial Practice]

Bid Form for the Procurement of Goods
[shall be submitted with the Bid]

BID FORM

Date : _____
 Project Identification No. : _____

To: *[name and address of Procuring Entity]*

Having examined the Philippine Bidding Documents (PBDs) including the Supplemental or Bid Bulletin Numbers *[insert numbers]*, the receipt of which is hereby duly acknowledged, we, the undersigned, offer to *[supply/deliver/perform]* *[description of the Goods]* in conformity with the said PBDs for the sum of *[total Bid amount in words and figures]* or the total calculated bid price, as evaluated and corrected for computational errors, and other bid modifications in accordance with the Price Schedules attached herewith and made part of this Bid. The total bid price includes the cost of all taxes, such as, but not limited to: *[specify the applicable taxes, e.g. (i) value added tax (VAT), (ii) income tax, (iii) local taxes, and (iv) other fiscal levies and duties]*, which are itemized herein or in the Price Schedules,

If our Bid is accepted, we undertake:

- a. to deliver the goods in accordance with the delivery schedule specified in the Schedule of Requirements of the Philippine Bidding Documents (PBDs);
- b. to provide a performance security in the form, amounts, and within the times prescribed in the PBDs;
- c. to abide by the Bid Validity Period specified in the PBDs and it shall remain binding upon us at any time before the expiration of that period.

[Insert this paragraph if Foreign-Assisted Project with the Development Partner:

Commissions or gratuities, if any, paid or to be paid by us to agents relating to this Bid, and to contract execution if we are awarded the contract, are listed below:

Name and address of agent	Amount and Purpose of Commission or gratuity
---------------------------	--

(if none, state “None”)]

Until a formal Contract is prepared and executed, this Bid, together with your written acceptance thereof and your Notice of Award, shall be binding upon us.

We understand that you are not bound to accept the Lowest Calculated Bid or any Bid you may receive.

We certify/confirm that we comply with the eligibility requirements pursuant to the PBDs.

The undersigned is authorized to submit the bid on behalf of *[name of the bidder]* as evidenced by the attached *[state the written authority]*.

We acknowledge that failure to sign each and every page of this Bid Form, including the attached Schedule of Prices, shall be a ground for the rejection of our bid.

Name: _____

Legal capacity: _____

Signature: _____

Duly authorized to sign the Bid for and behalf of: _____

Date: _____

Price Schedule for Goods Offered from Within the Philippines
[shall be submitted with the Bid if bidder is offering goods from within the Philippines]

For Goods Offered from Within the Philippines

Name of Bidder _____ Project ID No. _____ Page ___ of ___

1	2	3	4	5	6	7	8	9	10
Item	Description	Country of origin	Quantity	Unit price EXW per item	Transportation and all other costs incidental to delivery, per item	Sales and other taxes payable if Contract is awarded, per item	Cost of Incidental Services, if applicable, per item	Total Price, per unit (col 5+6+7+8)	Total Price delivered Final Destination (col 9) x (col 4)

Name: _____

Legal Capacity: _____

Signature: _____

Duly authorized to sign the Bid for and behalf of: _____

Price Schedule for Goods Offered from Abroad
[shall be submitted with the Bid if bidder is offering goods from Abroad]

For Goods Offered from Abroad

Name of Bidder _____ Project ID No. _____ Page ___ of ___

1	2	3	4	5	6	7	8	9
Item	Description	Country of origin	Quantity	Unit price CIF port of entry (specify port) or CIP named place (specify border point or place of destination)	Total CIF or CIP price per item (col. 4 x 5)	Unit Price Delivered Duty Unpaid (DDU)	Unit price Delivered Duty Paid (DDP)	Total Price delivered DDP (col 4 x 8)

Name: _____

Legal Capacity: _____

Signature: _____

Duly authorized to sign the Bid for and behalf of: _____



PHILIPPINE HEART CENTER
East Avenue, Quezon City

Abstract of Sealed Bidding re: Pharmaceutical Supplies for the period of January 1, 2022 to December 31,2022

BID FORM

S P E C I F I C A T I O N S	Average Monthly Consumption	AGENCY UNIT COST	BID PRICE	BIDDER
A. Medicines Acting on the Nervous System				
ANALGESICS/ANTIPYRETICS				
Opioid Analgesics				
1	Butorphanol			
	.01 2mg/ml, 1ml ampule	5	454.02	
2	Tramadol hydrochloride			
	.01 50mg capsule	2,000	1.98	
	.02 50mg/ml, 2ml ampule	1,000	6.38	
Non-Opioid Analgesics				
3	ASA (Acetyl Salicylic Acid)			
	.01 Aspirin 300mg tablet	1,000	2.60	
	.02 Aspirin 100mg Enteric Coated tablet	6,000	2.75	
	.03 Aspirin 80mg tablet	100	0.65	
	.04 Aspirin 80mg Enteric Coated tablet	33,600	2.73	
4	Ketorolac			
	.01 30mg/ml, 1ml ampule	400	14.42	
5	Paracetamol			
	.01 500mg tablet	15,000	0.45	
	.02 250mg/5ml, 60ml syrup OR suspension	144	17.25	
	.03 100mg/ml, 15ml Oral Drops (alcohol-free)	40	15.00	
	.04 150mg/ml, 2ml ampule	2,000	4.12	
	.05 10mg/ml, 100ml vial	2,000	107.33	
	.06 10mg/ml, 50ml vial	500	198.00	
ANTI-CONVULSANTS				
6	Carbamazepine			
	.01 200mg tablet	200	1.69	
7	Divalproex sodium			
	.01 250mg Extended Release tablet	200	8.88	
	.02 500mg Extended Release tablet	200	11.38	
8	Gabapentin			
	.01 300mg capsule	100	6.25	
	.02 100mg capsule	100	5.60	
9	Levetiracetam			
	.01 500mg Film Coated tablet	600	16.38	
	.02 100mg/ml, 5ml vial	200	1,734.00	
10	Phenobarbital			
	.01 Phenobarbital 60mg tablet	100	5.09	
	.02 Phenobarbital 30mg tablet	100	3.80	
	.03 Luminal 130mg/ml ampule	100	489.79	
11	Phenytoin			
	.01 100mg capsule	4,000	11.50	
	.02 50mg/ml, 5ml ampule	10	595.00	
	.03 50mg/ml, 2ml ampule	10	83.28	
12	Valproic acid			
	.01 500mg/5ml vial	150	1,694.65	
	.02 250mg/5ml, 120ml syrup	10	314.05	

S P E C I F I C A T I O N S		Average Monthly Consumption	AGENCY UNIT COST	BID PRICE	BIDDER
13	ANTI-DEPRESSANTS				
	Sertraline .01 50mg tablet	300	7.25		
14	ANTI-PAKINSON DRUG				
	Carbidopa + Levodopa .01 25mg + 100mg tablet	50	14.55		
15	ANTI-DEMENTIA				
	Memantine hydrochloride .01 10mg tablet	200	18.00		
16	CEREBRAL STIMULANTS				
	Mecobalamin .01 500mcg tablet	200	11.00		
17	ANESTHETIC AGENTS				
	Bupivacaine .01 0.5%, 10ml Duofit polyamp (green)	40	93.04		
18	.02 0.5%, 5ml ampule (heavy)	3	92.33		
	Isoflurane + Vaporizer (with quarterly calibration) .01 100ml vial	12	649.44		
19	Lidocaine hydrochloride .01 2% 20mg/ml 5ml polyamp	3,000	10.00		
	.02 10%/dose, 50ml Spray	10	1,993.91		
20	Propofol .01 10mg/ml, 20ml ampule or vial	1,400	56.38		
	.02 10mg/ml, 50ml vial	200	189.63		
21	Sevoflurane (300-1000 ppm) + vaporizer .01 250ml PEN bottle	60	4,433.11		
22	PARASYMPATHOMIMETIC AGENTS				
	Neostigmine .01 0.5mg/1ml ampule	80	118.75		
23	DANGEROUS DRUGS/NARCOTICS				
	Ephedrine sulfate .01 50mg/ml ampule	150	81.50		
24	Fentanyl citrate .01 50mcg/ml, 10ml ampule	700	247.00		
	.02 50mcg/ml, 2ml ampule	1,500	46.78		
25	Ketamine hydrochloride .01 50mg/ml, 10ml vial	20	498.77		
26	Morphine sulfate .01 10mg/ml, 1ml ampule	300	49.70		
	.02 MST 10mg MR tablet	30	20.00		
27	Nalbuphine hydrochloride .01 10mg/ml ampule	400	44.88		
28	Oxycodone hydrochloride .01 10mg/ml, 1ml ampule	100	618.69		
	.02 10mg/ml, 2ml ampule	50	923.50		
	.03 10mg capsule	56	127.16		
29	Pethidine hydrochloride .01 50mg/ml, 2ml ampule	30	148.00		
30	Sedatives-Benzodiazepine				
	Alprazolam .01 500mcg tablet	600	15.75		
31	Bromazepam .01 1.5mg tablet	500	8.65		
32	Clonazepam .01 2mg tablet	1,500	3.00		

S P E C I F I C A T I O N S		Average Monthly Consumption	AGENCY UNIT COST	BID PRICE	BIDDER
33	Diazepam				
	.01 5mg tablet	1,000	9.29		
	.02 5mg/ml, 2ml ampule	300	74.34		
34	Midazolam				
	.01 15mg tablet	100	24.68		
	.02 5mg/ml, 1ml ampule	2,000	61.40		
	.03 1mg/ml, 5ml ampule	1,000	70.00		
	Non-Benzodiazepine				
35	Zolpidem				
	.01 10mg tablet	90	38.50		
	Benzodiazepine poisoning antidote				
36	Flumazenil				
	.01 100mcg/ml, 5ml apule	20			
B. Medicines Acting on the Musculoskeletal System and Joints					
	ANTI-GOUT PREPARATIONS				
37	Allopurinol				
	.01 100mg tablet	1,000	1.61		
	.02 300mg tablet	1,000	2.00		
38	Colchicine				
	.01 0.5mg tablet	1,200	2.10		
	ANTI-INFLAMMATORY (Non-Steroidal Drugs)				
39	Celecoxib				
	.01 200mg capsule	2,100	3.40		
	.02 400mg capsule	300	13.00		
40	Diclofenac sodium				
	.01 1% Emulgel, 20gram tube	3	378.75		
41	Ibuprofen				
	.01 100mg/5ml, 60ml suspension	20	37.76		
42	Mefenamic Acid				
	.01 250mg tablet	200	1.07		
	.02 500mg tablet	1,200	0.91		
	.03 500mg Super Fast (SF) capsule	800	36.25		
	SKELETAL MUSCLE RELAXANTS				
43	Atracurium besylate				
	.01 10mg/ml, 2.5ml ampule	300	77.84		
44	Rocuronium bromide				
	.01 10mg/ml, 5ml vial	1,200	158.60		
45	Sugammadex				
	.01 100mg/ml, 2ml vial	40	5,257.07		
C. Anti-Infectives					
	AMIBESIDE				
46	Metronidazole				
	.01 500mg tablet	300	1.20		
	.02 125mg/5ml, 60ml suspension	5	20.25		
	.03 5mg/ml, 100ml vial	400	14.17		
	AMINOGLYCOSIDE				
47	Amikacin sulfate				
	.01 250mg/ml, 2ml ampule	1,000	43.33		
48	Gentamicin sulfate				
	.01 40mg/ml, 2ml ampule	300	3.40		
49	Streptomycin				
	.01 1 gram vial	10	21.20		
	CEPHALOSPORINS				
	1st GENERATION				
50	Cefalexin				
	.01 500mg capsule	300	2.65		
	.02 250mg/5ml, 70ml suspension	10	25.93		

S P E C I F I C A T I O N S		Average Monthly Consumption	AGENCY UNIT COST	BID PRICE	BIDDER
51	Cefazolin				
	.01 1 gram vial	100	16.53		
52	2nd GENERATION Cefoxitin (Close Drug Delivery System)				
	.01 1 gram Solution for Infusion in DUPLEX, bag	480	814.87		
53	Cefuroxime				
	.01 750mg vial	3,000	16.30		
	.02 1.5 gram vial (alternate)				
	.03 500mg tablet	2,000	9.00		
	.04 250mg/5ml, 120ml suspension	30	199.00		
54	3rd GENERATION Cefixime				
	.01 200mg capsule	800	7.74		
55	Ceftazidime				
	.01 1 gram vial	600	34.00		
56	Ceftriaxone + 10ml diluent for Intravenous Injection				
	.01 1 gram vial	2,000	18.56		
57	4th GENERATION Cefipime				
	.01 1 gram vial	1,440	84.38		
58	MACROLIDES Azithromycin dihydrate				
	.01 500mg tablet	960	8.98		
	.02 500mg vial	60	322.00		
	.03 200mg/5ml, 15ml suspension	10	230.00		
59	Clarithromycin				
	.01 500mg tablet	300	10.64		
	.02 500mg OD tablet	90	189.00		
60	Erythromycin				
	.01 250mg tablet	30	8.84		
61	PENICILLINS Ampicillin				
	.01 500mg vial	500	8.21		
62	Ampicillin + Sulbactam				
	.01 500mg + 250mg vial	600	21.14		
	.02 1gram + 500mg vial	600	58.17		
63	Amoxicillin trihydrate				
	.01 500mg capsule	2,000	1.28		
	.02 250mg/5ml 60ml suspension	40	20.48		
	.03 100mg/ml, 15ml Oral drops	3	19.80		
64	Amoxicillin + Clavulanic acid				
	.01 500mg + 125mg tablet	2,000	7.00		
	.02 875mg + 125mg tablet	420	10.99		
	.03 200mg + 28.5mg/5ml, 70ml suspension	10	115.76		
	.04 400mg + 57mg/5ml, 70ml suspension	10	168.37		
65	Benzathine Benzyl Penicillin				
	.01 1.2 million units vial	200	49.55		
66	Cloxacillin sodium				
	.01 500mg capsule	300	2.78		
67	Oxacillin				
	.01 500mg vial	600	17.88		

S P E C I F I C A T I O N S		Average Monthly Consumption	AGENCY UNIT COST	BID PRICE	BIDDER
68	Penicillin G sodium				
	.01 Benzyl PCN 1 million units vial	60	6.24		
	.02 Benzyl PCN 5 million units vial	60	15.92		
69	Phenoxymethyl Penicillin V potassium				
	.01 500mg capsule	1,000	15.44		
	.02 250mg capsule	15,000	9.00		
70	Piperacillin sodium + Tazobactam sodium				
	.01 2 grams + 250mg vial	2,000	72.83		
	.02 4 grams + 500mg vial	2,400	97.44		
	CARBAPENEM				
71	Ertapenem				
	.01 1 gram vial	150	2,562.57		
72	Meropenem trihydrate				
	.01 1 gram vial	1,000	156.45		
	.02 500mg vial	600	104.39		
	TETRACYCLINES				
73	Doxycycline hydrochloride				
	.01 100mg capsule	200	1.23		
	OTHER ANTIBIOTICS				
74	Amphotericin B				
	.01 50mg Lyophilized Powder for Injection (non-lipid)	30	988.00		
75	Aztreonam				
	.01 1 gram vial	180	1,244.00		
76	Clindamycin				
	.01 150mg capsule	300	2.75		
	.02 300mg capsule	1,000	5.35		
	.03 150mg/ml, 4ml ampule	800	73.00		
77	Colistimethate sodium (Colistin)				
	.01 2 million IU vial	300	1,730.00		
78	Linezolid				
	.01 600mg tablet	30	1,790.23		
	.02 300mg/ml, 2ml solution for injection bag	60	2,525.00		
79	Polymyxin B sulfate				
	.01 500,000 Units, 5ml vial	300	2,386.00		
80	Vancomycin hydrochloride				
	.01 500mg vial	2,000	137.22		
	Sulfa Derivatives				
81	Cotrimoxazole (Trimethoprim+Sulfamethoxazole)				
	.01 160mg + 800mg tablet	200	1.90		
	.02 200mg + 40mg / 5ml, 60ml suspension	4	15.40		
	QUINOLONES				
82	Ciprofloxacin				
	.01 500mg tablet	2,000	1.30		
	.02 100mg Extended Release tablet	28	61.72		
	.03 2mg/ml, 100ml vial	200	23.78		
	.04 2mg/ml, 200ml vial	200	1,187.00		
83	Levofloxacin				
	.01 500mg tablet	400	6.13		
	.02 750 mg tablet	200	13.50		
	.03 5mg/ml, 100ml vial	600	104.22		
	.04 5mg/ml, 150ml vial	300	1,356.00		

S P E C I F I C A T I O N S		Average Monthly Consumption	AGENCY UNIT COST	BID PRICE	BIDDER
ANTI-TUBERCULOSIS DRUGS					
84	Isoniazid + Vitamin B6				
	.01 200mg + 12mg / 5ml, 120ml syrup	3	70.00		
85	Rifampicin				
	.01 200mg/5ml, 100ml syrup	3	158.17		
	.02 200mg/5ml, 120ml syrup	3	265.00		
86	Rifampicin + Isoniazid				
	.01 150mg + 75mg tablet	200	6.00		
87	Rifampicin + Ethambutol + Isoniazid + Pyrazinamide				
	.01 150mg + 275mg + 75mg + 400mg tablet	2,400	6.49		
ANTI-VIRAL / ANTI-FUNGAL					
88	Acyclovir				
	.01 400mg tablet	150	11.15		
89	Fluconazole				
	.01 2mg/ml, 100ml vial	160	312.34		
	.02 150mg capsule	40	68.00		
	.03 50mg capsule	560	69.40		
90	Nystatin				
	.01 100,000 units/ml, 30ml oral suspension	30	155.00		
D. Immunologicals					
SERUM, TOXOIDS/VACCINES/ANTI-VIRAL					
91	Hepatitis-B Vaccine				
	.01 20mcg / ml, 1ml vial	20	165.75		
92	Immunoglobulin, IV				
	.01 10%, 2 grams, 20ml vial (alternate)				
	.02 5%, 2.5 grams, 50ml vial	100	4,260.00		
93	Influenza Vaccine (trivalent or tetravalent)				
	.01 0.5ml prefilled syringe	500	550.00		
94	Pneumococcal vaccine				
	.01 25mcg / 0.5ml prefilled syringe	5	2,561.47		
95	Tetanus Antitoxin				
	.01 1,500 units vial	10	100.00		
96	Tetanus Toxoid				
	.01 0.5ml ampule	20	33.68		
97	Tetanus Immune Globulin (Human)				
	.01 250 IU, 1ml prefilled syringe	20	615.00		
E. Cardiovascular Medicines					
Inotropics					
98	Adenosine				
	.01 3mg/ml, 2ml vial or ampule	120	221.31		
99	Digoxin				
	.01 0.25mg tablet	12,000	3.97		
	.02 0.05mg/ml, 60ml Elixir	70	567.00		
	.03 0.25mg/ml, 2ml ampule	300	108.34		
100	Dobutamine hydrochloride				
	.01 50mg/ml, 5ml ampule	1,500	145.85		
101	Dopamine hydrochloride				
	.01 40mg/ml, 5ml ampule	400	37.45		
102	Epinephrine hydrochloride				
	.01 1mg/ml, 1ml ampule	2,400	21.91		
103	Norepinephrine bitartrate				
	.01 1mg/ml, 4ml ampule	2,000	148.04		
	.02 1mg/ml, 10ml ampule	2,000	515.20		

S P E C I F I C A T I O N S		Average Monthly Consumption	AGENCY UNIT COST	BID PRICE	BIDDER
104	ANTI-ARRHYTHMICS				
	Amiodarone				
	.01 200mg tablet	3,000	16.28		
	.02 50mg/ml, 3ml ampule	1,000	200.00		
105	ANTI-PLATELET AGGREGANTS				
	Cilostazol				
	.01 50mg tablet	6,000	7.72		
	.02 100mg tablet	3,000	12.85		
106	Clopidogrel				
	.01 75mg Film Coated tablet	53,760	1.11		
	.02 75mg tablet (for post-OP use)	1,400	67.50		
107	BETA BLOCKERS				
	Atenolol				
	.01 50mg tablet	3,000	1.79		
	.02 100mg tablet	300	2.81		
108	Bisoprolol				
	.01 5mg tablet	2,000	13.50		
109	Carvedilol				
	.01 25mg tablet	24,000	4.27		
	.02 6.25mg tablet	84,000	1.18		
110	Esmolol				
	.01 10mg/ml, 10ml vial	200	499.00		
111	Metoprolol tartrate				
	.01 50mg tablet	20,000	0.95		
	.02 100mg tablet	10,000	1.55		
112	Propranolol				
	.01 10mg tablet	3,000	5.50		
	.02 40mg tablet	600	8.45		
113	ACE INHIBITOR				
	Captopril				
	.01 25mg tablet	3,000	0.80		
114	Enalapril maleate				
	.01 5mg tablet	10,000	2.86		
	.02 20mg tablet	3,000	7.25		
115	ANGIOTENSIN 2 RECEPTOR BLOCKERS				
	Irbesartan				
	.01 150mg tablet	9,000	3.12		
	.02 300mg tablet	3,360	7.98		
116	Irbesartan + Hydrochlorothiazide				
	.01 150mg + 12.5mg tablet	840	7.98		
117	Losartan potassium				
	.01 50mg tablet	75,600	0.80		
	.02 100mg. tablet	9,000	2.50		
118	Losartan + Hydrochlorothiazide				
	.01 50mg + 12.5mg tablet	9,000	3.01		
119	Sacubitril+Valsartan				
	.01 50mg tablet	11200	55.25		
	.02 100mg tablet	11200	56.25		
	.03 200mg tablet	6048	56.25		
120	Telmisartan				
	.01 40mg tablet	8,400	13.62		
	.02 80mg tablet	3,600	25.00		
121	Telmisartan + Hydrochlorothiazide				
	.01 40mg + 12.5mg tablet	3,600	19.02		

S P E C I F I C A T I O N S		Average Monthly Consumption	AGENCY UNIT COST	BID PRICE	BIDDER
122	Valsartan				
	.01 80mg tablet	1,800	8.42		
	.02 160mg tablet	300	11.00		
123	Valsartan + Hydrochlorothiazide				
	.01 80mg + 12.5mg tablet	560	11.49		
ALPHA-ADRENERGIC CENTRAL STIMULANTS					
124	Clonidine				
	.01 75mcg tablet	2,000	4.95		
	.02 150mcg tablet	1,000	11.00		
CALCIUM BLOCKERS					
125	Amlodipine besilate				
	.01 5mg tablet	120,000	0.39		
	.02 10mg tablet	20,000	0.53		
126	Diltiazem				
	.01 60mg tablet	4,000	12.00		
127	Felodipine				
	.01 5mg Extended or Monitored Release tablet	4,000	7.20		
	.02 10mg Extended or Monitored Release tablet	600	11.00		
128	Nicardepine hydrochloride				
	.01 1mg/ml, 10ml ampule	2,000	190.12		
129	Nifedipine				
	.01 30mg GITS tablet	300	41.25		
130	Nimodipine				
	.01 30mg tablet	500	27.52		
131	Verapamil hydrochloride				
	.01 80mg tablet	800	18.75		
	.02 240mg Sustained Release tablet	600	19.68		
	.03 2.5mg/ml, 2ml ampule	60	112.40		
PARASYMPATHOLYTIC AGENT					
132	Atropine sulfate				
	.01 1mg/ml, 1ml ampule	500	10.38		
ANTI-ANGINA					
133	Isosorbide dinitrate				
	.01 5mg Sublingual tablet	2,000	8.25		
	.02 10mg Oral tablet	600	9.00		
	.03 0.1%, 1mg/ml, 10ml ampule	1,200	131.78		
134	Isosorbide - 5 Mononitrate				
	.01 60mg Sustained Release capsule/tablet	6,000	7.00		
	.02 60mg Film Coated tablet	1,000	7.70		
	.03 30mg Sustained Release capsule	10,000	7.65		
	.04 30mg Film Coated tablet	1,000	11.10		
135	Nitroglycerine / Glyceryl trinitate				
	.01 1mg/ml, 10ml ampule	2,000	400.00		
136	Trimetazidine				
	.01 35mg Monitored Release tablet	84,000	3.77		
F. Diuretics					
137	Bumetamide				
	.01 1mg tablet	200	23.20		
	.02 0.5mg/ml 4ml ampule	60	77.75		

S P E C I F I C A T I O N S		Average Monthly Consumption	AGENCY UNIT COST	BID PRICE	BIDDER
138	Furosemide				
	.01 40mg tablet	14,000	1.50		
	.02 20mg tablet	8,000	1.00		
	.03 10mg/ml, 2ml ampule	12,000	5.43		
139	Hydrochlorothiazide				
	.01 25mg tablet	1,400	2.95		
140	Spirolactone				
	.01 25mg tablet	14,000	9.20		
	.02 50mg tablet	2,000	24.96		
141	Tolvaptan				
	.01 15mg tablet	200	652.80		
HYPOLIPIDEMIC AGENTS					
Fibric Acid Derivatives					
142	Fenofibrate				
	.01 200 Micronized capsule	2,400	10.87		
	.02 160mg tablet	1,500	27.50		
HMG Coenzyme A Reductase Inhibitors					
143	Atorvastatin calcium				
	.01 10mg tablet	3,000	4.86		
	.02 20mg tablet	12,000	4.98		
	.03 40mg tablet	24,000	9.24		
	.04 80mg tablet	16,000	19.18		
144	Rosuvastatin calcium				
	.01 10mg tablet	10,000	4.03		
	.02 20mg tablet	10,000	6.48		
145	Simvastatin				
	.01 20mg tablet	5,000	1.40		
	.02 40mg tablet	2,000	3.50		
G. Respiratory Medicines					
ANTITUSSIVES/Decongestants/Other Cough Remedies					
146	Butamirate citrate				
	.01 50mg Sustained Release tablet	1,000	12.50		
	.02 7.5mg/5ml 120ml syrup	6	120.00		
147	Dextromethorphan hydrobromide				
	.01 5mg Lozenges (6 loz per pack)	2	68.00		
BRONCHODILATORS					
148	Aminophylline				
	.01 25mg/ml, 10ml ampule	200	19.23		
149	Budesonide				
	.01 250mcg/ml, 2ml nebulizing solution	2,000	35.00		
150	Budesonide + Formoterol				
	.01 160mcg+4.5mcg turbuhaler, 60 doses	120	834.44		
151	Fluticasone propionate				
	.01 250mcg/ml, 2ml nebule	10	69.00		
152	Fluticasone + Formoterol				
	.01 250mcg/10mcg metered dose inhaler (MDI)	3	1,859.16		
153	Indacaterol+Glycopyrronium				
	.01 110mcg/50mcg capsule	300	62.48		
154	Ipratropium bromide				
	.01 0.5mg/2ml, 2ml Unit Dose Vial (UDV)	400	79.11		
155	Ipratropium + Salbutamol				
	.01 500mcg+2.5mg/2.5ml pulmoneb	10,000	9.65		

S P E C I F I C A T I O N S		Average Monthly Consumption	AGENCY UNIT COST	BID PRICE	BIDDER
156	Montelukast				
	.01 10 mg tablet	900	7.25		
157	Salbultamol				
	.01 100mcg/dose , 200 doses Metred Dose Inhaler	200	76.84		
	.02 2mg/5ml, 60ml syrup	5	14.98		
	.03 2.5mg/2.5ml nebule	10,000	8.00		
158	Salmeterol + Fluticasone				
	.01 500 diskus, 60 doses	10	463.98		
	.02 250 diskus, 60 doses	20	428.00		
	.03 100 diskus, 60 doses	3	389.85		
	.04 25mcg+250mcg inhaler, 120 doses	60	283.38		
	.05 25mcg+125mcg inhaler, 120 doses	60	222.10		
159	Tiotropium bromide				
	.01 18mcg/dose capsule	150	57.33		
	.02 Handihaler	5	305.00		
EXPECTORANTS / MUCOLYTICS					
160	Acetylcysteine				
	.01 600mg Effervescent tablet	12,000	18.90		
	.02 200mg granule/sachet	600	8.50		
	.03 100mg granule/sachet	450	8.00		
	.04 100mg/ml, 3ml (IM)ampule for injection	500	107.93		
161	Carbocysteine				
	.01 500mg capsule	600	10.50		
H. Antiallergics					
ANTIHISTAMINES					
162	Cetirizine				
	.01 10mg tablet	4,000	0.40		
	.02 5mg/5ml, 30ml syrup	60	60.00		
163	Diphenhydramine hydrochloride				
	.01 25mg capsule	2,000	0.75		
	.02 50mg/ml, 1ml ampule	600	18.78		
	.03 12.5mg/5 ml, 60 ml syrup	10	18.66		
164	Loratadine				
	.01 10mg tablet	500	1.47		
	.02 5mg/5ml, 30ml syrup	4	89.00		
I. Medicines Affecting the Blood					
ANTI-COAGULANT					
Unfractionated Heparin					
165	Heparin sodium				
	.01 1000 IU/ml, 5ml vial	5,000	46.49		
	.02 5000 IU/ml, 5ml vial	1,000	148.44		
Low Molecular Weight Heparin					
166	Enoxaparine sodium				
	.01 20mg/0.2ml Prefilled syringe	100	471.60		
	.02 40mg/0.4ml Prefilled syringe	3,000	195.60		
	.03 60mg/0.6ml Prefilled syringe	3,000	278.32		
167	Fondaparinux sodium				
	.01 2.5 mg/0.5ml Prefilled syringe	150	1,100.00		
168	Tinzaparin sodium				
	.01 10,000 IU/2ml vial	5	796.96		
	.02 10,000 IU/ml, 0.35 ml Prefilled syringe	60	432.62		

S P E C I F I C A T I O N S		Average Monthly Consumption	AGENCY UNIT COST	BID PRICE	BIDDER
ORAL ANTICOAGULANT					
169	Warfarin sodium				
	.01 1mg tablet	1,000	15.43		
	.02 2.5mg tablet	12,000	12.00		
	.03 5mg tablet	12,000	12.00		
HEMOSTATICS					
170	Epoetin-alfa				
	.01 4000 IU/0.4ml Prefilled syringe	1,000	350.00		
171	Epoetin-beta				
	.01 5000 IU/0.3ml Prefilled syringe	400	978.00		
172	Tranexamic Acid				
	.01 500mg capsule	300	5.38		
	.02 100mg/ml, 5ml ampule	3,000	14.32		
173	Vitamin K1/Phytomenadione				
	.01 10mg/ml, 1ml ampule	400	16.77		
IRON PREPARATIONS					
174	Ferrous sulfate (equiv to 60mg elemental iron)				
	.01 325mg tablet	4,000	1.00		
175	Ferrous gluconate +Vitamin B12+ Vitamin C				
	.01 (equivalent to Fe 30mg) capsule	1,000	24.00		
176	Ferrous sulfate + Multivitamins				
	.01 149.34mg/5ml, 120ml syrup	72	286.75		
	.02 74.64mg/ml, 15ml drops	5	46.78		
177	Ferrous fumarate/sulfate + Multivitamins + Folic acid				
	.01 tablet	1,000	0.69		
178	Folic acid				
	.01 5mg capsule or tablet	1,600	3.10		
PLASMINOGEN ACTIVATORS/THROMBOLYTICS					
179	Alteplase / recombinant-Tissue Plasminogen Activator				
	.01 50mg, 50ml vial	6	27,264.30		
180	Streptokinase				
	.01 1.5 million units vial	2	3,700.00		
J. Blood Products and Blood Substitute					
BLOOD DERIVATIVES					
181	Albumin, human (with transfusion set)				
	.01 25%, 50ml vial	1,000	2,400.00		
	.02 20%, 50ml vial	300	1,648.00		
PLASMA EXPANDER					
182	Hydroxyethyl starch				
	.01 6%, 500ml bag	10	380.00		
183	Modified Fluid Gelatin				
	.01 4%, 500ml bottle	200	515.00		
K. Gastrointestinal Medicines					
ANTACIDS / H2ANTAGONISTS/ACID-PUMP INHIBITOR					
184	Famotidine				
	.01 20mg FC tablet	300	22.00		
	.02 20mg vial	300	88.33		
185	Lanzoprazole				
	.01 30mg FDT tablet	56	75.74		
186	Magnesium+Aluminum hydroxide				
	.01 180ml suspension	40	297.00		
187	Omeprazole				
	.01 20mg capsule	1,200	0.81		
	.02 40mg capsule	8,000	4.23		
	.03 40mg vial	3,000	24.78		

S P E C I F I C A T I O N S		Average Monthly Consumption	AGENCY UNIT COST	BID PRICE	BIDDER
188	Pantoprazole				
	.01 20mg EC tablet	420	30.00		
	.02 40mg EC tablet	1,400	40.00		
189	Rabeprazole				
	.01 20mg tablet	140	13.00		
190	Sucralfate				
	.01 1 Gram tablet	400	30.50		
ANTI-DIARRHEALS					
191	Loperamide				
	.01 2mg capsule	300	0.92		
192	Rifaximin				
	.01 200mg tablet	300	63.20		
ANTI-EMETIC / ANTI-VERTIGO					
193	Betahistine				
	.01 16mg tablet	600	11.00		
	.02 24mg tablet	600	30.00		
194	Cinnarizine				
	.01 25mg tablet	200	1.20		
195	Domperidone				
	.01 10mg tablet	1,000	2.00		
196	Metoclopramide				
	.01 10mg tablet	200	1.73		
	.02 5mg/ml, 2ml ampule	600	3.97		
ANTI-FLATULENCE / ANTI-SPASMODIC / BILE SALT					
197	Hyoscine n-butylbromide				
	.01 10mg tablet	600	3.84		
	.02 20mg/ml ampule	100	17.23		
198	Ursodeoxycholic acid				
	.01 250mg capsule	300	34.14		
LAXATIVES / CATHARTICS					
199	Bisacodyl				
	.01 5mg tablet	600	1.78		
	.02 5mg suppository (Pediatric)	50	1.54		
	.03 10mg suppository (Adult)	300	19.43		
200	Castor Oil				
	.01 60ml bottle	20	35.00		
201	Lactulose				
	.01 3.3grams/5ml, 120ml syrup	600	92.50		
202	Monobasic + Dibasic Sodium Phosphate				
	.01 19Grams + 7Grams / 118ml (Adult)	20	205.00		
	.02 48Grams+18Grams /100ml, 45ml solution (flavored)	10	271.00		
L. Hormone and Hormone Antagonists					
CORTICOSTEROIDS					
203	Dexamethasone				
	.01 4mg tablet	300	18.00		
	.02 5mg/ml ampule	2,000	23.62		
204	Hydrocortisone vial + diluent				
	.01 100mg dry pack + 2ml diluent vial	2,000	18.78		
205	Methylprednisolone				
	.01 4mg tablet	200	5.00		
	.02 16mg tablet	300	8.20		
	.03 40mg/ml, 1ml vial	100	289.95		
	.04 500mg vial	20	1,155.00		

S P E C I F I C A T I O N S		Average Monthly Consumption	AGENCY UNIT COST	BID PRICE	BIDDER
206	Prednisone				
	.01 5mg tablet	600	0.59		
	.02 20mg tablet	600	2.36		
HYPOLYCEMIC AGENTS (ORAL)					
207	Gliclazide				
	.01 30mg OD tablet	3,000	2.60		
	.02 60mg Monitored Release tablet	6,000	9.28		
	.03 80mg tablet	600	3.12		
208	Metformin				
	.01 500mg tablet	24,000	0.54		
	.02 500mg Extended Release tablet	6,000	7.58		
	.03 850mg tablet	200	4.39		
I N S U L I N S					
Rapid Acting and Short Duration					
209	Regular				
	.01 100 IU/ml, 10ml vial	400	92.00		
	.02 100 IU/ml, 5ml vial	20	73.00		
Intermediate and Long Acting					
210	NPH (Isophane)				
	.01 100 IU/ml, 10ml vial	480	79.00		
Long Acting					
211	Biphasic Isophane 70% + Recombinant DNA 30%				
	.01 100 IU/ml, 10ml vial	2,000	89.00		
THYROID/ ANTI-THYROID PREPARATIONS					
212	Levothyroxin sodium				
	.01 50mcg tablet	2,000	3.44		
	.02 100mcg tablet	2,000	6.47		
213	Methimazole				
	.01 5mg tablet	1,000	5.41		
214	Propyl-thiouracil (5 PTU)				
	.01 50mg tablet	600	5.52		
M. Medicines Correcting Water Electrolyte / Parenteral Nutrition					
ELECTROLYTES / ELEMENTS					
215	Calcium carbonate				
	.01 500mg capsule or tablet	3,000	1.65		
216	Calcium + Vitamin D				
	.01 600mg+200 IU Film-Coated tablet	3,000	4.00		
217	Calcium gluconate				
	.01 10%, 10ml ampule	6,000	15.11		
218	Hypertonic lactate solution				
	.01 250ml Flexi bag	90	717.70		
219	Magnesium sulfate				
	.01 25%, 10ml ampule	3,000	23.44		
220	Oral Rehydration Salt / Electrolyte				
	.01 Chloride 65meq+Sodium 75meq Oral Solution, 500ml	3	115.00		
221	Potassium citrate				
	.01 10 meq tablet	600	6.80		
222	Potassium chloride				
	.01 750mg Durule tablet	6,000	12.00		
	.02 2 meq/ml, 20ml vial	1,600	23.00		
223	Sodium bicarbonate				
	.01 650mg tablet	1,000	0.99		
	.02 8.4%, 50ml vial	2,000	90.00		

S P E C I F I C A T I O N S		Average Monthly Consumption	AGENCY UNIT COST	BID PRICE	BIDDER
224	Sodium chloride				
	.01 1 Gram tablet	600	6.25		
	.02 2.5meq/ml, 20ml vial	200	19.00		
	.03 0.9% Sodium chloride 10ml polyamp	16,000	11.00		
	PARENTAL NUTRITION				
225	Amino Acids, Essential				
	.01 6% 100ml bottle for infant	200	359.00		
	.02 8%, 500ml bag	30	506.00		
	.03 10%, 500ml bottle	5	730.00		
226	Dextrose 50% IV				
	.01 50%, 50ml vial	2,000	19.60		
227	Glucose + Amino Acids + Lipids				
	.01 Peripheral 1400 kilocalorie bag	40	1,525.00		
	.02 Lipid 1500 kilocalorie bag	40	3,500.00		
	INTRAVENOUS FLUIDS				
228	.01 Dextrose 5% in Water, 250ml (Glass)	300	54.49		
	.02 Dextrose 5% in Water, 500ml	1,200	27.22		
	.03 Dextrose 10% in Water, 500ml	144	26.32		
	.04 Dextrose 5% in 0.3% Sodium Chloride, 500ml (Glass)	1,200	34.98		
	.05 Dextrose 5% in 0.3% Sodium Chloride, 500ml	1,680	29.52		
	.06 Dextrose 5% in 0.9% Sodium Chloride, 500ml	240	27.72		
	.07 Dextrose 5% in 0.9% Sodium Chloride, 1000ml	360	29.96		
	.08 Dextrose 5% in Lactated Ringer's Soln, 500ml	72	32.36		
	.09 Dextrose 5% in Lactated Ringer's Soln, 1000ml	144	30.00		
	.10 Isotonic Electrolyte Solution, 1000ml	600	120.00		
	.11 Ionosol in Dextrose 5% in Water, 500ml	144	36.00		
	.12 Lactated Ringer's Solution, 500ml	2,400	32.70		
	.13 Lactated Ringer's Solution, 1000ml (alternate)				
	.14 0.9% Sodium Chloride, 50ml	100	21.11		
	.15 0.9% Sodium Chloride, 100ml	300	49.00		
	.16 0.9% Sodium Chloride, 250ml bag	60	99.00		
	.17 0.9% Sodium Chloride, 500ml bag	60	101.00		
	.18 0.9% Sodium Chloride, 500ml	12,000	28.89		
	.19 0.9% Sodium Chloride, 1000ml bag	1,000	138.00		
	.20 0.9% Sodium Chloride, 1000ml	6,000	26.60		
	.21 Balanced Multiple Maintenance Solution with 5% Dextrose, 1000ml.	120	38.90		
	.22 Balanced Multiple Replacement Solution with 5% Dextrose, 500ml.	120	28.99		
	.23 Balanced Multiple Replacement Solution with 5% Dextrose, 1000ml.	600	38.90		

S P E C I F I C A T I O N S		Average Monthly Consumption	AGENCY UNIT COST	BID PRICE	BIDDER
229	Mannitol				
	.01 20%, 500ml bottle	360	69.38		
230	Essential Fatty Acids + Phospholipids + Vitamin E				
	.01 MCT / LCT 10%, 500ml bottle	5	600.00		
231	Sterile Water for Injection				
	.01 Sterile water for injection 1000ml bottle (twist-off)	1,200	34.00		
	.02 Sterile water for injection 1000ml bottle (glass)	24	104.50		
	.03 Sterile water for injection 500ml bottle (glass)	150	105.00		
	.04 Sterile water for injection, 100ml	100	44.73		
	.05 Sterile water for injection, 50ml	1,440	20.17		
	.06 Sterile water for injection, 10ml	9,600	10.50		
N. Diagnostic Agents					
	Contrast Media for: Kidneys/Blood Vessels	c/o	Diagnostic Lab.		
232	Gadobutrol				
	.01 5 ml vial	60	2,550.00		
233	Gadodiamide				
	.01 287mg/ml vial, 10ml	5	2,500.00		
234	Gadoteric acid				
	.01 279.32mg/ml vial	10	2,700.00		
235	Iopamidol				
	.01 300mg, 50ml	30	712.80		
	.02 370mg, 50ml	480	1,002.50		
	.03 370mg, 100ml	400	1,450.80		
236	Iopromide				
	.01 300mg vial, 50ml	30	808.50		
	.02 370mg vial, 50ml	160	1,475.00		
	.03 370mg vial, 100ml	200	1,900.00		
237	Iohexol				
	.01 300mg Iodine/ml vial, 50ml	60	988.48		
	.02 350mg Iodine/ml vial, 50ml	150	980.98		
238	Ioversol 741mg/ml				
	.01 300, 50ml	60	1,200.00		
	.02 350, 50ml	60	1,300.00		
O. Dermatologicals and Ophthalmological Preparations					
	DERMAL ANTIBIOTICS				
239	Acyclovir				
	.01 5%, 2 Gram cream	5	724.00		
240	Fusidate sodium				
	.01 5 Gram ointment	5	163.34		
241	Mupirocin				
	.01 2% 5 Grams cream (alternate)				
	.02 2% 5 Grams ointment	400	44.33		
242	Silver sulfadiazine				
	.01 10mg/Gram, 20 Gram cream	200	69.11		
	DERMAL ANTIFUNGAL				
243	Ketoconazole				
	.01 2%, 5 Gram cream	3	215.50		
244	Terbinafine				
	.01 1%, 10 Gram cream	5	465.25		
	DERMAL STEROID				
245	Betamethasone dipropionate				
	.01 0.05%, 5 Gram cream	3	41.74		

S P E C I F I C A T I O N S		Average Monthly Consumption	AGENCY UNIT COST	BID PRICE	BIDDER
246	Fluocinolone acetonide				
	.01 0.25%, 5 Gram cream	3	310.00		
P. Throat Preparations					
247	Povidone iodine				
	.01 1% Oromucosal solution, 120ml bottle (gargle)	576	153.47		
Q. Vitamins and Minerals / Food Supplements					
248	Multivitamins+Minerals				
	.01 capsule or tablet	9,600	1.93		
	.02 60ml or 100ml or 120ml syrup	144	44.00		
	.03 15ml oral drops	20	22.85		
249	Multivitamins+Zinc				
	.01 capsule or film-coated tablet	10,000	10.00		
	.02 120ml syrup	144	44.00		
250	Vitamin - B Complex				
	.01 Vitamin B1 (500mg) tablet	10,000	18.00		
	.02 Vitamin B6 (200mg-250mg) tablet	4,000	11.00		
	.03 Vitamin B12 (5mg) tablet	300	48.25		
251	Vitamin C (Ascorbic Acid)				
	.01 500mg tablet	25,000	0.80		
	.02 100mg/5ml, 120ml syrup	216	28.00		
	.03 100mg/ml, 15ml oral drops	60	17.50		
R. Chemotherapeutic Medicines / Antineoplastics					
252	Calcium folinate				
	.01 10mg/ml, 5ml vial	3	155.00		
253	Capecitabine				
	.01 500mg tablet	360	37.00		
254	Carboplatin				
	.01 10mg/ml, 15ml vial	3	887.00		
	.02 10mg/ml, 45ml vial	5	1,600.00		
255	Cisplatin				
	.01 1mg/ml, 10ml vial	5	160.00		
	.02 1mg/ml, 50ml vial	5	366.17		
256	Cyclophosphamide				
	.01 1 gram vial	5	210.00		
257	Docetaxel				
	.01 10mg/ml, 8ml vial	5	2,486.50		
	.02 10mg/ml, 2ml vial	5	586.00		
258	Doxorubicin				
	.01 2mg/ml, 5ml vial	3	151.00		
259	Epirubicin				
	.01 2mg/ml, 25ml vial	3	1,600.00		
	.02 2mg/ml, 5ml vial	3	637.00		
260	Filgrastim (GCSF)				
	.01 300mcg/ml prefilled syringe	3	1,115.00		
261	Gemcitabine				
	.01 1 gram vial	5	1,700.00		
	.02 200mg vial	5	499.00		
262	Paclitaxel				
	.01 100mg vial	5	980.00		
	.02 30mg vial	5	488.00		
263	Trastuzumab				
	.01 150mg vial	3	11,895.95		