

PHILIPPINE BIDDING DOCUMENTS

**PROCUREMENT OF
DRUGS AND MEDICINE
ITB # 2015-003
March 27, 2015
9:00 a.m.**

BATANGAS MEDICAL CENTER

**Third Edition
October 2009**

Preface

These Philippine Bidding Documents (PBDs) for the procurement of Goods through Competitive Bidding have been prepared by the Government of the Philippines (GOP) for use by all branches, agencies, departments, bureaus, offices, or instrumentalities of the government, including government-owned and/or –controlled corporations (GOCCs), government financial institutions (GFIs), state universities and colleges (SUCs), and local government units (LGUs). The procedures and practices presented in this document have been developed through broad experience, and are for mandatory¹ use in projects that are financed in whole or in part by the GOP or any foreign government/foreign or international financing institution in accordance with the provisions of the Implementing Rules and Regulations (IRR) of Republic Act (RA) 9184.

The Bidding Documents shall clearly and adequately define, among others: (a) the objectives, scope, and expected outputs and/or results of the proposed contract; (b) the eligibility requirements of bidders, such as track record to be determined by the Head of the Procuring Entity; (c) the expected contract duration, the estimated quantity in the case of procurement of goods, delivery schedule and/or time frame; and (d) the obligations, duties, and/or functions of the winning bidder.

In order to simplify the preparation of the Bidding Documents for each procurement, the PBDs groups the provisions that are intended to be used unchanged in

Section II. Instructions to Bidders (ITB) and in

Section IV. General Conditions of Contract (GCC). Data and provisions specific to each procurement and contract should be included in Section III. Bid Data Sheet (BDS);

¹ Unless the Treaty or International or Executive Agreement expressly provides use of foreign government/foreign or international financing institution procurement procedures and guidelines.

Section V. Special Conditions of Contract (SCC); Section VI. Schedule of Requirements; and

Section VII. Technical Specifications. The forms to be used are provided in where the information is useful for the Bidder. 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 Project.
- (b) Specific details, such as the name of the Procuring Entity and address for bid submission, should be furnished in the ITB, BDS, and SCC. The final documents should contain neither blank spaces nor options.

This Preface and the footnotes or notes in italics included in the Invitation to Bid, BDS, SCC, 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. The Bidding Documents should contain no footnotes except

since these provide important guidance to Bidders.

- (c) The cover should be modified as required to identify the Bidding Documents as to the names of the Project, Contract, and Procuring Entity, in addition to date of issue.
- (d) If modifications must be made to bidding procedures, they can be presented in the BDS. Modifications for specific Project or Contract details should be provided in the SCC as amendments to the Conditions of Contract. For easy completion, whenever reference has to be made to specific clauses in the BDS or SCC these terms shall be printed in bold type face on Section I. Instructions to Bidders and Section III. General Conditions of Contract, respectively.

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Republic of the Philippines
Department of Health CHD IV
BATANGAS MEDICAL CENTER
(formerly Batangas Regional Hospital)
Batangas City

INVITATION TO BID

The Batangas Medical Center through the General Appropriations Act, now invite bids for the Delivery of the Following:

Particulars	Approved Budget for the Contract	Cost of Bid Docs	Pre-bid Conference	Opening of Bid
Security Services	2,700,000.00	5,000.00	Nov. 28, 2013 9:30 a.m.	Dec. 17, 2013 9:30 a.m.
X-ray supplies	6,000,000.00	25,000.00	Nov. 28, 2013 9:30 a.m.	Dec. 17, 2013 1:30 p.m.
Medical oxygen	9,400,000.00	10,000.00	Nov. 28, 2013 9:30 a.m.	Dec. 18, 2013 9:30 a.m.
Laboratory Supplies & rgt	24,000,000.00	25,000.00	Nov. 28, 2013 9:30 a.m.	Dec. 18, 2013 1:30 p.m.

Drugs & medicines	28,000,000.00	10,000.00	Nov. 28, 2013 9:30 a.m.	Dec. 19, 2013 9:30 a.m.
Med/surgical supplies	31,000,000.00	25,000.00	Nov. 28, 2013 9:30 a.m.	Dec. 23, 2013 9:30 a.m.
Dorm & CHD-IV-A extension office	10,000,000.00	10,000.00	Nov. 28, 2013 9:30 a.m.	Dec. 27, 2013 9:30 a.m.
Rehabilitation of TB wards & Isolation rooms	3,800,000.00	5,000.00	Nov. 28, 2013 9:30 a.m.	Dec. 27, 2013 1:30 p.m.

Bids received in excess of the Approved Budget Contract for each particulars shall be automatically rejected at bid opening. Bidders should have completed, within 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, Instruction to Bidders.

Bidding is restricted to Filipino citizen/ sole proprietorships, partnerships, or organization with at least sixty percent (60%) of outstanding capital stock belonging to citizens of the Philippines, and to citizens or organizations of a country, the laws or regulations or which grant similar right or privileges to Filipino citizens, pursuant to RA5183 and subject to Commonwealth Act 138.

Interested bidders may obtain further information from BatMC and inspect the bidding documents at Supply Office during Mondays to Fridays, 8:00 a.m. to 5:00 p.m except on holidays. A complete set of bidding documents may be purchased by interested bidders from November 25, 2013 up to December 17, 2013 from the address specified below, upon payment of a non-refundable fee for the bidding documents in the amount stated in the above table. The BatMC will hold a Pre-bid Conference on November 28, 2013, 9:30 a.m. which shall be open only to all interested parties who have purchased the Bidding Documents. All bids must be accompanied by a bid security in any of the acceptable forms and in the amount stated in ITB Clause 18. Bids will be opened in the presence of the Bidder's representatives who choose to attend the address below. Late bids shall not be accepted.

The Batangas Medical Center reserves the right to accept or reject any Bid, to annul the bidding process, to reject all Bids and may not award the contract without incurring any liability and make no assurance that a contract shall be entered into as a result the bidding when the funds for the particulars has been withheld or reduced though no fault of the BatMC at any time prior to contract award, without thereby incurring any liability to the affected Bidder or Bidders.

For more information, please refer to:

HBAC Secretarial
Batangas Medical Center
Batangas City
Telefax No. (043) 723-6176

ELIZABETH V. PALINES MD, FPNA
HBAC Chairperson

Section II. Instructions to Bidders

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A. General

1. Scope of Bid

The procuring entity named in the BDS (hereinafter referred to as the “Procuring Entity”) wishes to receive bids for supply and delivery of the goods as described in

1.1. Section VII. Technical Specifications (hereinafter referred to as the “Goods”).

1.2. The name, identification, and number of lots specific to this bidding are provided in the BDS. The contracting strategy and basis of evaluation of lots is described in ITB Clause 28.

2. Source of Funds

The Procuring Entity has a budget or has applied for or received funds from the Funding Source named in the BDS, and in the amount indicated in the BDS. It intends to apply part of the funds received for the Project, as defined in the BDS, to cover eligible payments under the contract.

3. Corrupt, Fraudulent, Collusive, and Coercive Practices

3.1. 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. In pursuance of this policy, the Procuring Entity:

(a) defines, for purposes of this provision, the terms set forth below as follows:

(i) “corrupt practice” means behavior on the part of officials in the public or private sectors by which they improperly and unlawfully enrich themselves, others, or induce others to do so, by misusing the position in which they are placed, and includes the offering, giving, receiving, or soliciting of anything of value to influence the action of any such official in the procurement process or in contract execution;

entering, on behalf of the government, into any contract or transaction manifestly and grossly disadvantageous to the same, whether or not the public officer profited or will profit thereby, and similar acts as provided in RA 3019.

- (ii) “fraudulent practice” means a misrepresentation of facts in order to influence a procurement process or the execution of a contract to the detriment of the Procuring Entity, and includes collusive practices among Bidders (prior to or after bid submission) designed to establish bid prices at artificial, non-competitive levels and to deprive the Procuring Entity of the benefits of free and open competition.
- (iii) “collusive practices” means a scheme or arrangement between two or more Bidders, with or without the knowledge of the Procuring Entity, designed to establish bid prices at artificial, non-competitive levels.
- (iv) “coercive practices” means harming or threatening to harm, directly or indirectly, persons, or their property to influence their participation in a procurement process, or affect the execution of a contract;
- (b) will reject a proposal for award if it determines that the Bidder recommended for award has engaged in any of the practices mentioned in this Clause for purposes of competing for the contract.

3.2. Further, the Procuring Entity will seek to impose the maximum civil, administrative, and/or criminal penalties available under applicable laws on individuals and organizations deemed to be involved in any of the practices mentioned in ITB Clause 3.1(a).

3.3. Furthermore, the Funding Source and the Procuring Entity reserve the right to inspect and audit records and accounts of a bidder or supplier in the bidding for and performance of a contract themselves or through independent auditors as reflected in the GCC Clause 3.

4. Conflict of Interest

4.1. All Bidders found to have conflicting interests shall be disqualified to participate in the procurement at hand, without prejudice to the imposition of appropriate administrative, civil, and criminal sanctions. A Bidder may be considered to have conflicting interests with another Bidder in any of the events described in paragraphs (a) through (c) below and a general conflict of interest in any of the circumstances set out in paragraphs (d) through (f) below:

- (a) A Bidder has controlling shareholders in common with another Bidder;

- (b) **A Bidder receives or has received any direct or indirect subsidy from any other Bidder;**
- (c) **A Bidder has the same legal representative as that of another Bidder for purposes of this bid;**
- (d) **A Bidder has a relationship, directly or through third parties, that puts them in a position to have access to information about or influence on the bid of another Bidder or influence the decisions of the Procuring Entity regarding this bidding process. This will include a firm or an organization who lends, or temporarily seconds, its personnel to firms or organizations which are engaged in consulting services for the preparation related to procurement for or implementation of the project if the personnel would be involved in any capacity on the same project;**
- (e) **A Bidder submits more than one bid in this bidding process. However, this does not limit the participation of subcontractors in more than one bid; or**
- (f) **A Bidder who participated as a consultant in the preparation of the design or technical specifications of the Goods and related services that are the subject of the bid.**

4.2. In accordance with Section 47 of the IRR of RA 9184, all Bidding Documents shall be accompanied by a sworn affidavit of the Bidder that it is not related to the Head of the Procuring Entity, members of the Bids and Awards Committee (BAC), members of the Technical Working Group (TWG), members of the BAC Secretariat, the head of the Project Management Office (PMO) or the end-user unit, and the project consultants, by consanguinity or affinity up to the third civil degree. On the part of the Bidder, this Clause shall apply to the following persons:

- (a) **If the Bidder is an individual or a sole proprietorship, to the Bidder himself;**
- (b) **If the Bidder is a partnership, to all its officers and members;**
- (c) **If the Bidder is a corporation, to all its officers, directors, and controlling stockholders; and**
- (d) **If the Bidder is a joint venture (JV), the provisions of items (a), (b), or (c) of this Clause shall correspondingly apply to each of the members of the said JV, as may be appropriate.**

Relationship of the nature described above or failure to comply with this Clause will result in the automatic disqualification of a Bidder.

5. Eligible Bidders

- 5.1. Unless otherwise indicated in the BDS, the following persons shall be eligible to participate in this bidding:
- (a) Duly licensed Filipino citizens/sole proprietorships;
 - (b) Partnerships duly organized under the laws of the Philippines and of which at least sixty percent (60%) of the interest belongs to citizens of the Philippines;
 - (c) Corporations duly organized under the laws of the Philippines, and of which at least sixty percent (60%) of the outstanding capital stock belongs to citizens of the Philippines;
 - (d) Cooperatives duly organized under the laws of the Philippines, and of which at least sixty percent (60%) of the interest belongs to citizens of the Philippines; and
 - (e) Persons/entities forming themselves into a JV, *i.e.*, a group of two (2) or more persons/entities that intend to be jointly and severally responsible or liable for a particular contract: Provided, however, that Filipino ownership or interest of the joint venture concerned shall be at least sixty percent (60%).
- 5.2. Foreign bidders may be eligible to participate when any of the following circumstances exist, as specified in the BDS:
- (a) When a Treaty or International or Executive Agreement as provided in Section 4 of the RA 9184 and its IRR allow foreign bidders to participate;
 - (b) Citizens, corporations, or associations of a country, included in the list issued by the GPPB, the laws or regulations of which grant reciprocal rights or privileges to citizens, corporations, or associations of the Philippines;
 - (c) When the Goods sought to be procured are not available from local suppliers; or
 - (d) When there is a need to prevent situations that defeat competition or restrain trade.
- 5.3. Government corporate entities may be eligible to participate only if they can establish that they (a) are legally and financially autonomous, (b) operate under commercial law, and (c) are not dependent agencies of the GOP or the Procuring Entity.
- 5.4. Unless otherwise provided in the BDS, the Bidder must have completed at least one contract similar to the Project the value of which, adjusted to current prices using the National Statistics Office consumer price index, must be at least equivalent to a percentage of the ABC stated in the BDS.

For this purpose, contracts similar to the Project shall be those described in the BDS, and completed within the relevant period stated in the Invitation to Bid and ITB Clause 12.1(a)(iii).

- 5.5. **Unless otherwise provided in the BDS, the Bidder must submit a computation of its Net Financial Contracting Capacity (NFCC) or a commitment from a Universal or Commercial Bank to extend a credit line in its favor if awarded the contract for this Project (CLC).**

The NFCC, computed using the following formula, must be at least equal to the ABC to be bid:

NFCC = [(Current assets minus current liabilities) (K)] minus the value of all outstanding or uncompleted portions of the projects under ongoing contracts, including awarded contracts yet to be started coinciding with the contract for this Project.

Where:

K = 10 for a contract duration of one year or less, 15 for a contract duration of more than one year up to two years, and 20 for a contract duration of more than two years.

The CLC must be at least equal to ten percent (10%) of the ABC for this Project. If issued by a foreign bank, it shall be confirmed or authenticated by a Universal or Commercial Bank. In the case of local government units (LGUs), the Bidder may also submit CLC from other banks certified by the *Bangko Sentral ng Pilipinas* (BSP) as authorized to issue such financial instrument.

6. Bidder's Responsibilities

- 6.1. The Bidder or its duly authorized representative shall submit a sworn statement in the form prescribed in Section VIII. Bidding Forms as required in ITB Clause 12.I (b)(iii)

6.2. The Bidder is responsible for the following:

- (a) **Having taken steps to carefully examine all of the Bidding Documents;**
- (b) **Having acknowledged all conditions, local or otherwise, affecting the implementation of the contract;**
- (c) **Having made an estimate of the facilities available and needed for the contract to be bid, if any;**
- (d) **Having complied with its responsibility to inquire or secure Supplemental/Bid Bulletin(s) as provided under ITB Clause 10.3.**
- (e) Ensuring that it is not "blacklisted" or barred from bidding by the GOP or any of its agencies, offices, corporations, or LGUs, including

foreign government/foreign or international financing institution whose blacklisting rules have been recognized by the GPPB;

- (f) Ensuring that 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;
- (g) Authorizing the Head of the Procuring Entity or its duly authorized representative/s to verify all the documents submitted;
- (h) Ensuring that the signatory is the duly authorized representative of the Bidder, and granted full power and authority to do, execute and perform any and all acts necessary and/or to represent the Bidder in the bidding, with the duly notarized Secretary's Certificate attesting to such fact, if the Bidder is a corporation, partnership, cooperative, or joint venture;
- (i) Complying with the disclosure provision under Section 47 of RA 9184 in relation to other provisions of RA 3019; and
- (j) Complying with existing labor laws and standards, in the case of procurement of services.

Failure to observe any of the above responsibilities shall be at the risk of the Bidder concerned.

- 6.3. **The Bidder is expected to examine all instructions, forms, terms, and specifications in the Bidding Documents. Unless otherwise indicated in the BDS, failure to furnish all information or documentation required in the Bidding Documents shall result in the rejection of the bid and the disqualification of the Bidder.**
- 6.4. **It shall be the sole responsibility of the Bidder to determine and to satisfy itself by such means as it considers necessary or desirable as to all matters pertaining to the contract to be bid, including: (a) the location and the nature of this Project; (b) climatic conditions; (c) transportation facilities; and (d) other factors that may affect the cost, duration, and execution or implementation of this Project.**
- 6.5. **The Procuring Entity shall not assume any responsibility regarding erroneous interpretations or conclusions by the prospective or eligible bidder out of the data furnished by the procuring entity.**

- 6.6. The Bidder shall bear all costs associated with the preparation and submission of his bid, and the Procuring Entity will in no case be responsible or liable for those costs, regardless of the conduct or outcome of the bidding process.
- 6.7. Before submitting their bids, the Bidder is deemed to have become familiar with all existing laws, decrees, ordinances, acts and regulations of the Philippines which may affect this Project in any way.
- 6.8. The Bidder should note that the Procuring Entity will accept bids only from those that have paid the nonrefundable fee for the Bidding Documents at the office indicated in the Invitation to Bid.

7. Origin of Goods

Unless otherwise indicated in the BDS, there is no restriction on the origin of goods other than those prohibited by a decision of the United Nations Security Council taken under Chapter VII of the Charter of the United Nations, subject to ITB Clause 27.1.

8. Subcontracts

- 8.1. Unless otherwise specified in the BDS, the Bidder may subcontract portions of the Goods to an extent as may be approved by the Procuring Entity and stated in the BDS. However, subcontracting of any portion shall not relieve the Bidder from any liability or obligation that may arise from the contract for this Project.
- 8.2. Subcontractors must comply with the eligibility criteria and the documentary requirements specified in the BDS. In the event that any subcontractor is found by the Procuring Entity to be ineligible, the subcontracting of such portion of the Goods shall be disallowed.
- 8.3. The Bidder may identify the subcontractor to whom a portion of the Goods will be subcontracted at any stage of the bidding process or during contract implementation. If the Bidder opts to disclose the name of the subcontractor during bid submission, the Bidder shall include the required documents as part of the technical component of its bid.

B. Contents of Bidding Documents

9. Pre-Bid Conference

- 9.1. If so specified in the BDS, a pre-bid conference shall be held at the venue and on the date indicated therein, to clarify and address the Bidders' questions on the technical and financial components of this Project.
- 9.2. Bidders are encouraged to attend the pre-bid conference to ensure that they fully understand the Procuring Entity's requirements. Non-attendance of the Bidder will in no way prejudice its bid; however, the

Bidder is expected to know the changes and/or amendments to the Bidding Documents discussed during the pre-bid conference.

- 9.3. Any statement made at the pre-bid conference shall not modify the terms of the Bidding Documents unless such statement is specifically identified in writing as an amendment thereto and issued as a Supplemental/Bid Bulletin.**

10. Clarification and Amendment of Bidding Documents

- 10.1. Bidders who have purchased the Bidding Documents may request for clarifications on any part of the Bidding Documents for an interpretation. Such a request must be in writing and submitted to the Procuring Entity at the address indicated in the BDS at least ten (10) calendar days before the deadline set for the submission and receipt of bids.**
- 10.2. Supplemental/Bid Bulletins may be issued upon the Procuring Entity's initiative for purposes of clarifying or modifying any provision of the Bidding Documents not later than seven (7) calendar days before the deadline for the submission and receipt of bids. Any modification to the Bidding Documents shall be identified as an amendment.**
- 10.3. Any Supplemental/Bid Bulletin issued by the BAC shall also be posted on the Philippine Government Electronic Procurement System (PhilGEPS) and the website of the Procuring Entity concerned, if available. It shall be the responsibility of all Bidders who secure the Bidding Documents to inquire and secure Supplemental/Bid Bulletins that may be issued by the BAC. However, Bidders who have submitted bids before the issuance of the Supplemental/Bid Bulletin must be informed and allowed to modify or withdraw their bids in accordance with ITB Clause 23.**

C. Preparation of Bids

11. Language of Bid

The bid, as well as all correspondence and documents relating to the bid exchanged by the Bidder and the Procuring Entity, shall be written in English. Supporting documents and printed literature furnished by the Bidder may be in another language provided they are accompanied by an accurate translation in English certified by the appropriate embassy or consulate in the Philippines, in which case the English translation shall govern for purposes of interpretation of the bid.

12. Documents Comprising the Bid: Eligibility and Technical Components

- 12.1. Unless otherwise indicated in the BDS, the first envelope shall contain the following eligibility and technical documents:**

(a) **Eligibility Documents –**

Class “A” Documents:

- (i) **Registration certificate from the Securities and Exchange Commission (SEC), Department of Trade and Industry (DTI) for sole proprietorship, or Cooperative Development Authority (CDA) for cooperatives, or any proof of such registration as stated in the BDS;**
- (ii) **Mayor’s permit issued by the city or municipality where the principal place of business of the prospective bidder is located;**
- (iii) **Statement of all its ongoing and completed government and private contracts within the period stated in the BDS, including contracts awarded but not yet started, if any. The statement shall include, for each contract, the following:**
 - (i) **name of the contract;**
 - (ii) **date of the contract;**
 - (iii) **kinds of Goods;**
 - (iv) **amount of contract and value of outstanding contracts;**
 - (v) **date of delivery; and**
 - (vi) **end user’s acceptance or official receipt(s) issued for the contract, if completed.**
- (iv) **Audited financial statements, stamped “received” by the Bureau of Internal Revenue (BIR) or its duly accredited and authorized institutions, for the preceding calendar year, which should not be earlier than two (2) years from bid submission;**
- (v) **NFCC computation or CLC in accordance with ITB Clause 5.5; and**

Class “B” Document:

- (vi) **If applicable, the 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.**
- (b) **Technical Documents –**

- (i) **Bid security in accordance with ITB Clause 18. If the Bidder opts to submit the bid security in the form of:**
 - (i) **a bank draft/guarantee or an irrevocable letter of credit issued by a foreign bank, it shall be accompanied by a confirmation from a Universal or Commercial Bank; or**
 - (ii) **a surety bond, it shall be accompanied by a certification by the Insurance Commission that the surety or insurance company is authorized to issue such instruments;**
- (ii) **Conformity with technical specifications, as enumerated and specified in Sections VI and VII of the Bidding Documents; and**
- (iii) **Sworn statement in accordance with Section 25.2(a)(iv) of the IRR of RA 9184 and using the form prescribed in Section VIII. Bidding Forms.**

13. Documents Comprising the Bid: Financial Component

13.1. Unless otherwise stated in the **BDS**, the financial component of the bid shall contain the following:

- (a) Financial Bid Form, which includes bid prices and the bill of quantities and the applicable Price Schedules, in accordance with ITB Clauses 0 and 15.4;
- (b) If the Bidder claims preference as a Domestic Bidder or Domestic Entity, a certification from the DTI, SEC, or CDA issued in accordance with ITB Clause 27; and
- (c) Any other document required in the **BDS**.

13.2 (a) Unless otherwise stated in the BDS, all bids that exceed the ABC shall not be accepted.

- (b) Unless otherwise indicated in the BDS, for foreign-funded procurement, a ceiling may be applied to bid prices provided the following conditions are met:
 - (i) Bidding Documents are obtainable free of charge on a freely accessible website. If payment of Bidding Documents is required by the procuring entity, payment could be made upon the submission of bids.
 - (ii) The procuring entity has procedures in place to ensure that the ABC is based on recent estimates made by the responsible unit of the procuring entity and that the estimates reflect the quality, supervision

and risk and inflationary factors, as well as prevailing market prices, associated with the types of works or goods to be procured.

- (iii) The procuring entity has trained cost estimators on estimating prices and analyzing bid variances.
- (iv) The procuring entity has established a system to monitor and report bid prices relative to ABC and engineer's/procuring entity's estimate.
- (v) The procuring entity has established a system to monitor and report bid prices relative to ABC and procuring entity's estimate. The procuring entity has established a monitoring and evaluation system of contract implementation to provide a feedback on actual total cost of goods and works.

14. Alternative Bids

Alternative Bids shall be rejected. For this purpose, alternative bid is an offer made by a Bidder in addition or as a substitute to its original bid which may be included as part of its original bid or submitted separately therewith for purposes of bidding. A bid with options is considered an alternative bid regardless of whether said bid proposal is contained in a single envelope or submitted in two (2) or more separate bid envelopes.

15. Bid Prices

15.1. The Bidder shall complete the appropriate Price Schedules included herein, stating the unit prices, total price per item, the total amount and the expected countries of origin of the Goods to be supplied under this Project.

15.2. The Bidder shall fill in rates and prices for all items of the Goods described in the Bill of Quantities. Bids not addressing or providing all of the required items in the Bidding Documents including, where applicable, Bill of Quantities, shall be considered non-responsive and, thus, automatically disqualified. In this regard, where a required item is provided, but no price is indicated, the same shall be considered as non-responsive, but specifying a "0" (zero) for the said item would mean that it is being offered for free to the Government.

15.3 The terms Ex Works (EXW), Cost, Insurance and Freight (CIF), Cost and Insurance Paid to (CIP), Delivered Duty Paid (DDP), and other trade terms used to describe the obligations of the parties, shall be governed by the rules prescribed in the current edition of the International Commercial Terms (INCOTERMS) published by the International Chamber of Commerce, Paris.

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

- (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), including all customs duties and sales and other taxes already paid or payable:

- (i.I) on the components and raw material used in the manufacture or assembly of Goods quoted ex works or ex factory; or

- (i.2) on the previously imported Goods of foreign origin quoted ex warehouse, ex showroom, or off-the-shelf and any Procuring Entity country sales and other taxes which will be payable on the Goods if the contract is awarded.

- (ii) The price for inland transportation, insurance, and other local costs incidental to delivery of the Goods to their final destination.

- (iii) The price of other (incidental) services, if any, listed in the **BDS**.

(b) For Goods offered from abroad:

- (i) Unless otherwise stated in the **BDS**, the price of the Goods shall be quoted 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, listed in the **BDS**.

15.5 Prices quoted by the Bidder shall be fixed during the Bidder's performance of the contract and not subject to variation or price escalation on any account, unless otherwise specified in the **BDS**. A bid submitted with an adjustable price quotation shall be treated as non-responsive and shall be rejected, pursuant to ITB Clause 24.

All bid prices shall be considered as fixed prices, and therefore not subject to price escalation during contract implementation, except under extraordinary circumstances as indicated in the **BDS** and specified in the GCC and its corresponding SCC provision.

16. Bid Currencies

16.1 Prices shall be quoted in the following currencies:

- (a) For Goods that the Bidder will supply from within the Philippines, the prices shall be quoted in Philippine Pesos.
- (b) For Goods that the Bidder will supply from outside the Philippines, the prices may be quoted in the currency(ies) stated in the **BDS**. 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.

16.2 If so allowed in accordance with ITB Clause 16.1, the Procuring Entity for purposes of bid evaluation and comparing the bid prices will convert the amounts in various currencies in which the bid price is expressed to Philippine Pesos at the foregoing exchange rates.

16.3 Unless otherwise specified in the **BDS**, payment of the contract price shall be made in Philippine Pesos.

17. Bid Validity

- i. **Bids shall remain valid for the period specified in the BDS which shall not exceed one hundred twenty (120) calendar days from the date of the opening of bids.**
- ii. **In exceptional circumstances, prior to the expiration of the Bid validity period, the Procuring Entity may request Bidders to extend the period of validity of their bids. The request and the responses shall be made in writing. The bid security described in ITB Clause 18 should also be extended corresponding to the extension of the bid validity period at the least. A Bidder may refuse the request without forfeiting its bid security, but his bid shall no longer be considered for further evaluation and award. A Bidder granting the request shall not be required or permitted to modify its bid.**

18. Bid Security

18.1. The bid security, issued in favor of the Procuring Entity, in the amount stated in the BDS shall be equal to the percentage of the ABC in accordance with the following schedule:

Form of Bid Security	Amount of Bid Security (Equal to Percentage of the ABC)
(a) Cash or cashier's/manager's check issued by a Universal or Commercial Bank.	Two percent (2%)

(b) Bank draft/guarantee or irrevocable letter of credit issued by a Universal or Commercial Bank: Provided, however, that it shall be confirmed or authenticated by a Universal or Commercial Bank, if issued by a foreign bank.	
(c) Surety bond callable upon demand issued by a surety or insurance company duly certified by the Insurance Commission as authorized to issue such security.	Five percent (5%)
(d) Any combination of the foregoing.	Proportionate to share of form with respect to total amount of security
(e) Bid Securing declaration	No percentage required

For biddings conducted by LGUs, the Bidder may also submit bid securities in the form of cashier's/manager's check, bank draft/guarantee, or irrevocable letter of credit from other banks certified by the BSP as authorized to issue such financial statement.

18.2. The bid security should be valid for the period specified in the **BDS**. Any bid not accompanied by an acceptable bid security shall be rejected by the Procuring Entity as non-responsive.

18.3. No bid securities shall be returned to bidders after the opening of bids and before contract signing, except to those that failed or declared as post-disqualified, upon submission of a written waiver of their right to file a motion for reconsideration and/or protest. Without prejudice on its forfeiture, bid securities shall be returned only after the bidder with the Lowest Calculated and Responsive Bid has signed the contract and furnished the performance security, but in no case later than the expiration of the bid security validity period indicated in ITB Clause 18.2.

18.4. Upon signing and execution of the contract pursuant to ITB Clause 32, and the posting of the performance security pursuant to ITB Clause 33, the successful Bidder's bid security will be discharged, but in no case later than the bid security validity period as indicated in the ITB Clause 18.2.

18.5. The bid security may be forfeited:

(a) if a Bidder:

- (i) withdraws its bid during the period of bid validity specified in ITB Clause 17;
- (ii) does not accept the correction of errors pursuant to ITB Clause (b);

- (iii) fails to submit the requirements within the prescribed period or a finding against their veracity as stated in ITB Clause 29.2; or
- (iv) submission of eligibility requirements containing false information or falsified documents;
- (v) submission of bids that contain false information or falsified documents, or the concealment of such information in the bids in order to influence the outcome of eligibility screening or any other stage of the public bidding;
- (vi) allowing the use of one's name, or using the name of another for purposes of public bidding;
- (vii) withdrawal of a bid, or refusal to accept an award, or enter into contract with the Government without justifiable cause, after the Bidder had been adjudged as having submitted the Lowest Calculated and Responsive Bid;
- (viii) refusal or failure to post the required performance security within the prescribed time;
- (ix) refusal to clarify or validate in writing its bid during post-qualification with a period of seven (7) calendar days from receipt of the request for clarification;
- (x) any document attempt by a bidder to unduly influence the outcome of the bidding in his favor;
- (xi) failure of the potential joint venture partners to enter into the joint venture after the bid is declared successful;
- (xii) all other acts that tend to defeat the purpose of the competitive bidding, such as habitually withdrawing from bidding, submitting late Bids or patently insufficient bid, for at least three (3) times within a year, except for valid reasons.

(b) If the successful Bidder:

19. Format and Signing of Bids

19.1. Bidders shall submit their bids through their duly authorized representative using the appropriate forms provided in

19.2. Forms as mentioned in ITB Clause 0 must be completed without any alterations to their format, and no substitute form shall be accepted. All blank spaces shall be filled in with the information requested.

19.3. The Bidder shall prepare and submit an original of the first and second envelopes as described in ITB Clauses 12 and 12.1(b)(iii). In the event of any discrepancy between the original and the copies, the original shall prevail.

19.4. The bid, except for unamended printed literature, shall be signed, and each and every page thereof shall be initialed, by the duly authorized representative/s of the Bidder.

19.5. Any interlineations, erasures, or overwriting shall be valid only if they are signed or initialed by the duly authorized representative/s of the Bidder.

20. Sealing and Marking of Bids

20.1. Unless otherwise indicated in the BDS, Bidders shall enclose their original eligibility and technical documents described in ITB Clause 12 in one sealed envelope marked “ORIGINAL - TECHNICAL COMPONENT”, and the original of their financial component in another sealed envelope marked “ORIGINAL - FINANCIAL COMPONENT”, sealing them all in an outer envelope marked “ORIGINAL BID”.

20.1. Each copy of the first and second envelopes shall be similarly sealed duly marking the inner envelopes as “COPY NO. ____ - TECHNICAL COMPONENT” and “COPY NO. ____ – FINANCIAL COMPONENT” and the outer envelope as “COPY NO. ____”, respectively. These envelopes containing the original and the copies shall then be enclosed in one single envelope.

20.3. The original and the number of copies of the Bid as indicated in the BDS shall be typed or written in indelible ink and shall be signed by the bidder or its duly authorized representative/s.

20.4 All envelopes shall:

(a) contain the name of the contract to be bid in capital letters;

(b) bear the name and address of the Bidder in capital letters;

(c) be addressed to the Procuring Entity’s BAC in accordance with ITB Clause 0;

(d) bear the specific identification of this bidding process indicated in the ITB Clause 1.2; and

(e) bear a warning “DO NOT OPEN BEFORE...” the date and time for the opening of bids, in accordance with ITB Clause 21.

20.5.If bids are not sealed and marked as required, the Procuring Entity will assume no responsibility for the misplacement or premature opening of the bid.

D. Submission and Opening of Bids

21. Deadline for Submission of Bids

Bids must be received by the Procuring Entity’s BAC at the address and on or before the date and time indicated in the **BDS**.

22. Late Bids

Any bid submitted after the deadline for submission and receipt of bids prescribed by the Procuring Entity, pursuant to ITB Clause 21, shall be declared “Late” and shall not be accepted by the Procuring Entity.

23. Modification and Withdrawal of Bids

23.1. The Bidder may modify its bid after it has been submitted; provided that the modification is received by the Procuring Entity prior to the deadline prescribed for submission and receipt of bids. The Bidder shall not be allowed to retrieve its original bid, but shall be allowed to submit another bid equally sealed, properly identified, linked to its original bid marked as “TECHNICAL MODIFICATION” or “FINANCIAL MODIFICATION” and stamped “received” by the BAC. Bid modifications received after the applicable deadline shall not be considered and shall be returned to the Bidder unopened.

23.2. A Bidder may, through a Letter of Withdrawal, withdraw its bid after it has been submitted, for valid and justifiable reason; provided that the Letter of Withdrawal is received by the Procuring Entity prior to the deadline prescribed for submission and receipt of bids.

23.3. Bids requested to be withdrawn in accordance with ITB Clause 23.1 shall be returned unopened to the Bidders. A Bidder may also express its intention not to participate in the bidding through a letter which should reach and be stamped by the BAC before the deadline for submission and receipt of bids. A Bidder that withdraws its bid shall not be permitted to submit another bid, directly or indirectly, for the same contract.

23.4. No bid may be modified after the deadline for submission of bids. No bid may be withdrawn in the interval between the deadline for submission of bids and the expiration of the period of bid validity specified by the Bidder on the Financial Bid Form. Withdrawal of a bid during this interval shall result in the forfeiture of the Bidder’s bid security, pursuant to ITB Clause 18.5, and the imposition of

administrative, civil and criminal sanctions as prescribed by RA 9184 and its IRR.

24. Opening and Preliminary Examination of Bids

24.1 shall be considered as “failed”. Otherwise, the BAC shall rate the said first bid envelope The BAC shall open the first bid envelopes of Bidders in public as specified in the **BDS** to determine each Bidder’s compliance with the documents prescribed in ITB Clause 12. For this purpose, the BAC shall check the submitted documents of each bidder against a checklist of required documents to ascertain if they are all present, using a non-discretionary “pass/fail” criterion. If a bidder submits the required document, it shall be rated “passed” for that particular **requirement. In this regard, bids that fail to include any requirement or are incomplete or patently insufficient as “passed”.**

24.1. Unless otherwise specified in the BDS, immediately after determining compliance with the requirements in the first envelope, the BAC shall forthwith open the second bid envelope of each remaining eligible bidder whose first bid envelope was rated “passed”. The second envelope of each complying bidder shall be opened within the same day. In case one or more of the requirements in the second envelope of a particular bid is missing, incomplete or patently insufficient, and/or if the submitted total bid price exceeds the ABC unless otherwise provided in ITB Clause **Error! Reference source not found.**, the BAC shall rate the bid concerned as “failed”. Only bids that are determined to contain all the bid requirements for both components shall be rated “passed” and shall immediately be considered for evaluation and comparison.

24.2. Letters of withdrawal shall be read out and recorded during bid opening, and the envelope containing the corresponding withdrawn bid shall be returned to the Bidder unopened. If the withdrawing Bidder’s representative is in attendance, the original bid and all copies thereof shall be returned to the representative during the bid opening. If the representative is not in attendance, the bid shall be returned unopened by registered mail. The Bidder may withdraw its bid prior to the deadline for the submission and receipt of bids, provided that the corresponding Letter of Withdrawal contains a valid authorization requesting for such withdrawal, subject to appropriate administrative sanctions.

24.3. If a Bidder has previously secured a certification from the Procuring Entity to the effect that it has previously submitted the above-enumerated Class “A” Documents, the said certification may be submitted in lieu of the requirements enumerated in ITB Clause 12.1(a), items (i) to (v).

24.4. In the case of an eligible foreign Bidder as described in ITB Clause 5, the Class “A” Documents described in ITB Clause 12.1(a) may be substituted with the appropriate equivalent documents, if any, issued by the country of the foreign Bidder concerned.

24.5. Each partner of a joint venture agreement shall likewise submit the requirements in ITB Clauses 12.1(a)(i) and 12.1(a)(ii). Submission of documents required under ITB Clauses 12.1(a)(iii) to 12.1(a)(v) by any of the joint venture partners constitutes compliance.

24.6. The Procuring Entity shall prepare the minutes of the proceedings of the bid opening that shall include, as a minimum: (a) names of Bidders, their bid price, bid security, findings of preliminary examination; and (b) attendance sheet. The BAC members shall sign the abstract of bids as read.

E. Evaluation and Comparison of Bids

25. Process to be Confidential

25.1. Members of the BAC, including its staff and personnel, as well as its Secretariat and TWG, are prohibited from making or accepting any kind of communication with any bidder regarding the evaluation of their bids until the issuance of the Notice of Award, unless otherwise allowed in the **BDS** or in the case of ITB Clause 26.

25.2. Any effort by a bidder to influence the Procuring Entity in the Procuring Entity's decision in respect of bid evaluation, bid comparison or contract award will result in the rejection of the Bidder's bid.

26. Clarification of Bids

To assist in the evaluation, comparison, and post-qualification of the bids, the Procuring Entity may ask in writing any Bidder for a clarification of its bid. All responses to requests for clarification shall be in writing. Any clarification submitted by a Bidder in respect to its bid and that is not in response to a request by the Procuring Entity shall not be considered.

27. Domestic Preference

27.1. Unless otherwise stated in the **BDS**, the Procuring Entity will grant a margin of preference for the purpose of comparison of bids in accordance with the following:

- a. The preference shall be applied when (i) the lowest Foreign Bid is lower than the lowest bid offered by a Domestic Bidder, or (ii) the lowest bid offered by a non-Philippine national is lower than the lowest bid offered by a Domestic Entity.
- b. For evaluation purposes, the lowest Foreign Bid or the bid offered by a non-Philippine national shall be increased by fifteen percent (15%).

- c. In the event that (i) the lowest bid offered by a Domestic Entity does not exceed the lowest Foreign Bid as increased, or (ii) the lowest bid offered by a non-Philippine national as increased, then the Procuring Entity shall award the contract to the Domestic Bidder/Entity at the amount of the lowest Foreign Bid or the bid offered by a non-Philippine national, as the case may be.
- d. If the Domestic Entity/Bidder refuses to accept the award of contract at the amount of the Foreign Bid or bid offered by a non-Philippine national within two (2) calendar days from receipt of written advice from the BAC, the Procuring Entity shall award to the bidder offering the Foreign Bid or the non-Philippine national, as the case may be, subject to post-qualification and submission of all the documentary requirements under these Bidding Documents.

27.2 A Bidder may be granted preference as a Domestic Entity subject to the certification from the DTI (in case of sole proprietorships), SEC (in case of partnerships and corporations), or CDA (in case of cooperatives) that the (a) sole proprietor is a citizen of the Philippines or the partnership, corporation, cooperative, or association is duly organized under the laws of the Philippines with at least seventy five percent (75%) of its interest or outstanding capital stock belonging to citizens of the Philippines, (b) habitually established in business and habitually engaged in the manufacture or sale of the merchandise covered by his bid, and (c) the business has been in existence for at least five (5) consecutive years prior to the advertisement and/or posting of the Invitation to Bid for this Project.

27.3 A Bidder may be granted preference as a Domestic Bidder subject to the certification from the DTI that the Bidder is offering unmanufactured articles, materials or supplies of the growth or production of the Philippines, or manufactured articles, materials, or supplies manufactured or to be manufactured in the Philippines substantially from articles, materials, or supplies of the growth, production, or manufacture, as the case may be, of the Philippines.

28. Detailed Evaluation and Comparison of Bids

28.1. The Procuring Entity will undertake the detailed evaluation and comparison of bids which have passed the opening and preliminary examination of bids, pursuant to ITB Clause 24, in order to determine the Lowest Calculated Bid.

28.2. The Lowest Calculated Bid shall be determined in two steps:

- a. The detailed evaluation of the financial component of the bids, to establish the correct calculated prices of the bids; and
- b. The ranking of the total bid prices as so calculated from the lowest to the highest. The bid with the lowest **price shall be** identified as the Lowest Calculated Bid.

28.3 The Procuring Entity's BAC shall immediately conduct a detailed evaluation of all bids rated "passed," using non-discretionary pass/fail criteria. Unless otherwise specified in the **BDS**, the BAC shall consider the following in the evaluation of bids:

- (a) Completeness of the bid. Unless the ITB specifically allows partial bids, bids not addressing or providing all of the required items in the Schedule of Requirements including, where applicable, bill of quantities, shall be considered non-responsive and, thus, automatically disqualified. In this regard, where a required item is provided, but no price is indicated, the same shall be considered as non-responsive, but specifying a "0" (zero) for the said item would mean that it is being offered for free to the Procuring Entity; and
- (b) Arithmetical corrections. Consider computational errors and omissions to enable proper comparison of all eligible bids. It may also consider bid modifications, if allowed in the **BDS**. Any adjustment shall be calculated in monetary terms to determine the calculated prices.

28.4. Based on the detailed evaluation of bids, those that comply with the above-mentioned requirements shall be ranked in the ascending order of their total calculated bid prices, as evaluated and corrected for computational errors, discounts and other modifications, to identify the Lowest Calculated Bid. Total calculated bid prices, as evaluated and corrected for computational errors, discounts and other modifications, which exceed the ABC shall not be considered.

28.5. Unless otherwise indicated in the **BDS**, the Procuring Entity's evaluation of bids shall only be based on the bid price quoted in the Financial Bid Form.

28.6. Bids shall be evaluated on an equal footing to ensure fair competition. For this purpose, all bidders shall be required to include in their bids the cost of all taxes, such as, but not limited to, value added tax (VAT), income tax, local taxes, and other fiscal levies and duties which shall be itemized in the bid form and reflected in the detailed estimates. Such bids, including said taxes, shall be the basis for bid evaluation and comparison.

29. Post-Qualification

29.1. The Procuring Entity shall determine to its satisfaction whether the Bidder that is evaluated as having submitted the Lowest Calculated Bid (LCB) complies with and is responsive to all the requirements and conditions specified in ITB Clauses 5, 12, and 12.1(b)(iii).

29.2. Within a non-extendible period of three (3) calendar days from receipt by the bidder of the notice from the BAC that it submitted the LCB, the Bidder shall submit the following documentary requirements:

- (a) Tax clearance per Executive Order 398, Series of 2005, as finally reviewed and approved by the BIR;
- (b) Latest income and business tax returns in the form specified in the BDS.
- (c) Certificate of Phil GEPS Registration; and
- (d) Other appropriate licenses and permits required by law and stated in the **BDS**.

Failure of the Bidder declared as Lowest Calculated Bid to duly submit the requirements under this Clause or a finding against the veracity of such shall be ground for forfeiture of the bid security and disqualification of the Bidder for award.

29.3. The determination shall be based upon an examination of the documentary evidence of the Bidder's qualifications submitted pursuant to ITB Clauses 12 and 12.1(b)(iii), as well as other information as the Procuring Entity deems necessary and appropriate, using a non-discretionary "pass/fail" criterion.

29.4. If the BAC determines that the Bidder with the Lowest Calculated Bid passes all the criteria for post-qualification, it shall declare the said bid as the Lowest Calculated Responsive Bid, and recommend to the Head of the Procuring Entity the award of contract to the said Bidder at its submitted price or its calculated bid price, whichever is lower.

29.5. A negative determination shall result in rejection of the Bidder's Bid, in which event the Procuring Entity shall proceed to the next Lowest Calculated

Bid to make a similar determination of that Bidder's capabilities to perform satisfactorily. If the second Bidder, however, fails the post qualification, the procedure for post qualification shall be repeated for the Bidder with the next Lowest Calculated Bid, and so on until the Lowest Calculated Responsive Bid is determined for contract award.

- 29.6. Within a period not exceeding seven (7) calendar days from the date of receipt of the recommendation of the BAC, the Head of the Procuring Entity shall approve or disapprove the said recommendation. In the case of GOCCs and GFIs, the period provided herein shall be fifteen (15) calendar days.

30. Reservation Clause

- 30.1. Notwithstanding the eligibility or post-qualification of a bidder, the Procuring Entity concerned reserves the right to review its qualifications at any stage of the procurement process if it has reasonable grounds to believe that a misrepresentation has been made by the said bidder, or that there has been a change in the Bidder's capability to undertake the project from the time it submitted its eligibility requirements. Should such review uncover any misrepresentation made in the eligibility and bidding requirements, statements or documents, or any changes in the situation of the Bidder which will affect its capability to undertake the project so that it fails the preset eligibility or bid evaluation criteria, the Procuring Entity shall consider the said Bidder as ineligible and shall disqualify it from submitting a bid or from obtaining an award or contract.

- 30.2. Based on the following grounds, the Procuring Entity reserves the right to reject any and all bids, declare a Failure of Bidding at any time prior to the contract award, or not to award the contract, without thereby incurring any liability, and make no assurance that a contract shall be entered into as a result of the bidding:

- (a) If there is *prima facie* evidence of collusion between appropriate public officers or employees of the Procuring Entity, or between the BAC and any of the bidders, or if the collusion is between or among the bidders themselves, or between a bidder and a third party, including any act which restricts, suppresses or nullifies or tends to restrict, suppress or nullify competition;
- (b) If the Procuring Entity's BAC is found to have failed in following the prescribed bidding procedures; or
- (c) For any justifiable and reasonable ground where the award of the contract will not redound to the benefit of the GOP as follows:
 - i) If the physical and economic conditions have significantly changed so as to render the project no longer economically, financially or technically feasible as determined by the head of the procuring entity;

ii) If the project is no longer necessary as determined by the head of the procuring entity; and

iii) If the source of funds for the project has been withheld or reduced through no fault of the Procuring Entity.

30.3. In addition, the Procuring Entity may likewise declare a failure of bidding when:

- (a) No bids are received;
- (b) All prospective bidders are declared ineligible;
- (c) All bids fail to comply with all the bid requirements or fail post-qualification; or
- (d) The bidder with the Lowest Calculated Responsive Bid (LCRB) refuses, without justifiable cause to accept the award of contract, and no award is made.

F. Award of Contract

31. Contract Award

31.1. Subject to ITB Clause 29, the Procuring Entity shall award the contract to the Bidder whose bid has been determined to be the LCRB.

31.2. Prior to the expiration of the period of bid validity, the Procuring Entity shall notify the successful Bidder in writing that its bid has been accepted, through a Notice of Award received personally or sent by registered mail or electronically, receipt of which must be confirmed in writing within two (2) days by the Bidder with the LCRB and submitted personally or sent by registered mail or electronically to the Procuring Entity.

31.3. Notwithstanding the issuance of the Notice of Award, award of contract shall be subject to the following conditions:

- (a) Submission of the valid JVA, if applicable, within ten (10) calendar days from receipt by the Bidder of the notice from the BAC that the Bidder has the LCRB;
- (b) Posting of the performance security in accordance with ITB Clause 33;
- (c) Signing of the contract as provided in ITB Clause 32; and
- (d) Approval by higher authority, if required.

- 31.4. At the time of contract award, the Procuring Entity shall not increase or decrease the quantity of goods originally specified in Section VI. Schedule of Requirements.

32. Signing of the Contract

- 32.1 At the same time as the Procuring Entity notifies the successful Bidder that its bid has been accepted, the Procuring Entity shall send the Contract Form to the Bidder, which contract has been provided in the Bidding Documents, incorporating therein all agreements between the parties.
- 32.1. Within ten (10) calendar days from receipt of the Notice of Award, the successful Bidder shall post the required performance security and sign and date the contract and return it to the Procuring Entity.
- 32.2. The Procuring Entity shall enter into contract with the successful Bidder within the same ten (10) calendar day period provided that all the documentary requirements are complied with.
- 32.3.** The following documents shall form part of the contract:
- (a) Contract Agreement;
 - (b) Bidding Documents;
 - (c) Winning bidder's bid, including the Technical and Financial Proposals, and all other documents/statements submitted;
 - (d) Performance Security;
 - (e) Credit line in accordance with ITB Clause 5.5, if applicable;
 - (f) Notice of Award of Contract; and
 - (g) Other contract documents that may be required by existing laws and/or specified in the BDS.

33. Performance Security

- 33.1. To guarantee the faithful performance by the winning Bidder of its obligations under the contract, it shall post a performance security within a maximum period of ten (10) calendar days from the receipt of the Notice of Award from the Procuring Entity and in no case later than the signing of the contract.
- 33.2 The performance security shall be denominated in Philippine Pesos and posted in favor of the Procuring Entity in an amount equal to the percentage of the total contract price in accordance with the following schedule:

Form of Performance Security	Amount of Performance Security (Equal to Percentage of the Total Contract Price)
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(a) Cash or cashier's/manager's check issued by a Universal or Commercial Bank.	Five percent (5%)
(b) Bank draft/guarantee or irrevocable letter of credit issued by a Universal or Commercial Bank: Provided, however, that it shall be confirmed or authenticated by a Universal or Commercial Bank, if issued by a foreign bank.	
(c) Surety bond callable upon demand issued by a surety or insurance company duly certified by the Insurance Commission as authorized to issue such security; and/or	Thirty percent (30%)
(d) Any combination of the foregoing.	Proportionate to share of form with respect to total amount of security

33.3 Failure of the successful Bidder to comply with the above-mentioned requirement shall constitute sufficient ground for the annulment of the award and forfeiture of the bid security, in which event the Procuring Entity shall initiate and complete the post qualification of the second Lowest Calculated Bid. The procedure shall be repeated until the LCRB is identified and selected for contract award. However if no Bidder passed post-qualification, the BAC shall declare the bidding a failure and conduct a re-bidding with re-advertisement.

34. Notice to Proceed

34.1 Within three (3) calendar days from the date of approval of the contract by the appropriate government approving authority, the Procuring Entity shall issue its Notice to Proceed to the Bidder.

34.2. The date of the Bidder's receipt of the Notice to Proceed will be regarded as the effective date of the contract, unless otherwise specified in the BDS.

35. Protest Mechanism

Decision of the procuring entity at any stage of the procurement process may be questioned in accordance with Section 55 of the Revised Implementing Rules and Regulations of Republic Act 9184.

Section III. Bid Data Sheet

Bid Data Sheet

ITB Clause	
0	The Procuring Entity is <i>Batangas Medical Center</i>
1.2	The lot(s) and reference is/are: <i>[insert name]</i>
2	The Funding Source is: The Government of the Philippines (GOP) in the amount of <i>SEVEN MILLION FIVE HUNDRED PESOS ONLY</i> The name of the Project is: <i>PROCUREMENT OF MEDICAL OXGEN/ COMPRESS AIR/ LIQUID OXYGEN</i>
5.2	<i>Select one, delete the other.</i> None of the circumstances mentioned in the ITB Clause exists in this Project. Foreign bidders, except those falling under ITB Clause 5.2(b), may not participate in this Project. <i>Or</i> Foreign bidders may participate in this Project in view of the following circumstance(s): <i>[State which of the circumstance(s) mentioned in the ITB Clause exists in the Project.]</i>
5.4	<i>Select one, delete the other.</i> “failure of bidding” or “monopoly that will defeat the purpose of public bidding”], the Bidder should comply with the following requirements: a) Completed at least two (2) similar contracts, the aggregate amount of which should be equivalent to at least <i>[State “fifty percent (50%)” in the case of Non-expendable Supplies or “twenty-five percent (25%)” in the case of Expendable Supplies]</i> of the ABC for this Project; and b) The largest of these similar contracts must be equivalent to at least half of the percentage of the ABC as required above. For this purpose, similar contracts shall refer to <i>[insert description of similar contracts or state “No further instructions”]</i> .
5.5	No further instructions.

6.3	No further instructions.
7	No further instructions.
8.1	“Subcontracting is not allowed.”
8.2	“Not applicable”.
9.1	The Procuring Entity will hold a pre-bid conference for this Project on <i>November 27, 2014 at 9:00 a.m. at BatMC Conference Room</i>
10.1	The Procuring Entity’s address is: <i>Batangas Medical Center</i> <i>Adelaida U. Untalan</i> <i>(043) 723-6176</i>
12.1	No further instructions.
12.1(a)(i)	“No other acceptable proof of registration is recognized.”
12.1(a)(iii)	The statement of all ongoing and completed government and private contracts shall include all such contracts within two (2) years prior to the deadline for the submission and receipt of bids.
13.1	“No additional Requirements”
	The ABC is <i>SEVEN MILLION FIVE HUNDRED PESOS ONLY</i> . Any bid with a financial component exceeding this amount shall not be accepted.
(iii)	“No incidental services are required.”
0	“No incidental services are required.”
15.5	Bid Prices shall be fixed. Adjustable price proposals shall be treated as non-responsive and shall be rejected.
0	Extraordinary circumstances refer to events that may be determined by the National Economic and Development Authority in accordance with the Civil Code of the Philippines, and upon the recommendation of the Procuring Entity
16(b)	The Bid prices for Goods supplied from outside of the Philippines shall be quoted in Philippine Pesos..
16.3	No further instructions.
17.i	Bids will be valid until <i>one year upon receipt of Notice of Award</i>

18.1	<p>The bid security shall be in the following amount:</p> <ol style="list-style-type: none"> 1. 2% of ABC, if bid security is in cash, cashier's/manager's check, bank draft/guarantee or irrevocable letter of credit; 2. 5% of ABC if bid security is in Surety Bond; or 3. Any combination of the foregoing proportionate to the share of form with respect to total amount of security.
18.2	The bid security shall be valid 120 days.
Error! Reference source not found.	<p><i>grounds for forfeiture of bid security:</i></p> <ol style="list-style-type: none"> 1. Submission of eligibility requirements containing false information or falsified documents. 2. Submission of bids that contain false information or falsified documents, or the concealment of such information in the bids in order to influence the outcome of eligibility screening or any other stage of the public bidding. 3. Allowing the use of one's name, or using the name of another for purposes of public bidding. 4. Withdrawal of a bid, or refusal to accept an award, or enter into contract with the Government without justifiable cause, after the Bidder had been adjudged as having submitted the Lowest Calculated and Responsive Bid. 5. Refusal or failure to post the required performance security within the prescribed time. 6. Refusal to clarify or validate in writing its bid during post-qualification within a period of seven (7) calendar days from receipt of the request for clarification. 7. Any documented unsolicited attempt by a bidder to unduly influence the outcome of the bidding in his favor. 8. Failure of the potential joint venture partners to enter into the joint venture after the bid is declared as successful. 9. All other acts that tend to defeat the purpose of the competitive bidding, such as habitually withdrawing from bidding, submitting late Bids or patently insufficient bid, for at least three (3) times within a year, except for valid reasons.
(iv)	No further instructions.

0	No further instructions.
0	Each Bidder shall submit <i>one (1)</i> original and <i>two (2)</i> copies of the first and second components of its bid.
21	The address for submission of bids is <i>Batangas Medical Center</i> . The deadline for submission of bids is <i>December 8, 2014, 9:00 a.m.</i>
24.1	The place of bid opening is <i>BatMC CONFERENCE ROOM</i> The date and time of bid opening is <i>December 8, 2014, 9:00 a.m.</i>
25.1	No further instructions.
27.1	“No further instructions. ”
28.3	<i>No further instructions</i>
(b)	<i>bid modification is allowed</i>
28.5	No further instructions.
	<i>tax returns filed through the Electronic Filing and Payments System (EFPS).</i>
Error! Reference source not found.	<i>Philgeps registration certificate</i>
34.2	The effective date of the Contract is <i>upon receipt of the Notice of Award/Proceed</i>

Section IV. General Conditions of Contract

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1. Definitions

1.1. In this Contract, the following terms shall be interpreted as indicated:

- (a) “The Contract” means the agreement entered into between the Procuring Entity and the Supplier, as recorded in the Contract Form signed by the parties, including all attachments and appendices thereto and all documents incorporated by reference therein.**
- (b) “The Contract Price” means the price payable to the Supplier under the Contract for the full and proper performance of its contractual obligations.**
- (c) “The Goods” means all of the supplies, equipment, machinery, spare parts, other materials and/or general support services which the Supplier is required to provide to the Procuring Entity under the Contract.**
- (d) “The Services” means those services ancillary to the supply of the Goods, such as transportation and insurance, and any other incidental services, such as installation, commissioning, provision of technical assistance, training, and other such obligations of the Supplier covered under the Contract.**
- (e) “GCC” means the General Conditions of Contract contained in this Section.**
- (f) “SCC” means the Special Conditions of Contract.**
- (g) “The Procuring Entity” means the organization purchasing the Goods, as named in the SCC.**

- (h) **“The Procuring Entity’s country” is the Philippines.**
- (i) **“The Supplier” means the individual contractor, manufacturer distributor, or firm supplying/manufacturing the Goods and Services under this Contract and named in the SCC.**
- (j) **The “Funding Source” means the organization named in the SCC.**
- (k) **“The Project Site,” where applicable, means the place or places named in the SCC.**
- (l) **“Day” means calendar day.**
- (m) **The “Effective Date” of the contract will be the date of receipt by the Supplier of the Notice to Proceed or the date provided in the Notice to Proceed. Performance of all obligations shall be reckoned from the Effective Date of the Contract.**
- (n) **“Verified Report” refers to the report submitted by the Implementing Unit to the Head of the Procuring Entity setting forth its findings as to the existence of grounds or causes for termination and explicitly stating its recommendation for the issuance of a Notice to Terminate.**

2. Corrupt, Fraudulent, Collusive, and Coercive Practices

2.1. The Procuring Entity as well as the bidders, contractors, or suppliers shall observe the highest standard of ethics during the procurement and execution of this Contract. In pursuance of this policy, the Procuring Entity:

- (a) **defines, for the purposes of this provision, the terms set forth below as follows:**
 - (i) **"corrupt practice" means behavior on the part of officials in the public or private sectors by which they improperly and unlawfully enrich themselves, others, or induce others to do so, by misusing the position in which they are placed, and it includes the offering, giving, receiving, or soliciting of anything of value to influence the action of any such official in the procurement process or in contract execution; entering, on behalf of the Government, into any contract or transaction manifestly and grossly disadvantageous to the same, whether or not the public officer profited or will profit thereby, and similar acts as provided in Republic Act 3019.**
 - (ii) **"fraudulent practice" means a misrepresentation of facts in order to influence a procurement process or the execution of a contract to the detriment of the Procuring Entity, and includes collusive practices among Bidders (prior to or after**

bid submission) designed to establish bid prices at artificial, non-competitive levels and to deprive the Procuring Entity of the benefits of free and open competition.

(iii) “collusive practices” means a scheme or arrangement between two or more Bidders, with or without the knowledge of the Procuring Entity, designed to establish bid prices at artificial, non-competitive levels.

(iv) “coercive practices” means harming or threatening to harm, directly or indirectly, persons, or their property to influence their participation in a procurement process, or affect the execution of a contract;

(b) will reject a proposal for award if it determines that the Bidder recommended for award has engaged in any of the practices mentioned in this Clause for purposes of competing for the contract.

2.2. Further the Funding Source, Borrower or Procuring Entity, as appropriate, will seek to impose the maximum civil, administrative and/or criminal penalties available under the applicable law on individuals and organizations deemed to be involved with any of the practices mentioned in GCC Clause 2.1(a).

3. Inspection and Audit by the Funding Source

The Supplier shall permit the Funding Source to inspect the Supplier’s accounts and records relating to the performance of the Supplier and to have them audited by auditors appointed by the Funding Source, if so required by the Funding Source.

4. Governing Law and Language

4.1. This Contract shall be interpreted in accordance with the laws of the Republic of the Philippines.

4.2. This Contract has been executed in the English language, which shall be the binding and controlling language for all matters relating to the meaning or interpretation of this Contract. All correspondence and other documents pertaining to this Contract exchanged by the parties shall be written in English.

5. Notices

5.1. Any notice, request, or consent required or permitted to be given or made pursuant to this Contract shall be in writing. Any such notice, request, or consent shall be deemed to have been given or made when received by the concerned party, either in person or through an authorized representative of the Party to whom the communication is addressed, or when sent by registered mail, telex, telegram, or facsimile to such Party at the address specified in the SCC, which shall be effective

when delivered and duly received or on the notice's effective date, whichever is later.

- 5.2. A Party may change its address for notice hereunder by giving the other Party notice of such change pursuant to the provisions listed in the SCC for GCC Clause 5.1.

6. Scope of Contract

- 6.1. The GOODS and Related Services to be provided shall be as specified in Section VI. Schedule of Requirements.

- 6.2. 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. Any additional requirements for the completion of this Contract shall be provided in the SCC.

7. Subcontracting

- 7.1. Subcontracting of any portion of the Goods, if allowed in the BDS, does not relieve the Supplier of any liability or obligation under this Contract. The Supplier will be responsible for the acts, defaults, and negligence of any subcontractor, its agents, servants or workmen as fully as if these were the Supplier's own acts, defaults, or negligence, or those of its agents, servants or workmen.

- 7.2. Subcontractors disclosed and identified during the bidding may be changed during the implementation of this Contract, subject to compliance with the required qualifications and the approval of the Procuring Entity.

8. Procuring Entity's Responsibilities

- 8.1. Whenever the performance of the obligations in this Contract requires that the Supplier obtain permits, approvals, import, and other licenses from local public authorities, the Procuring Entity shall, if so needed by the Supplier, make its best effort to assist the Supplier in complying with such requirements in a timely and expeditious manner.

- 8.2. The Procuring Entity shall pay all costs involved in the performance of its responsibilities in accordance with GCC Clause 6.

9. Prices

Prices charged by the Supplier for Goods delivered and/or services performed under this Contract shall not vary from the prices quoted by the Supplier in its bid, with the exception of any change in price resulting from a Change Order issued in accordance with GCC Clause 29, or if applicable, adjustments authorized in accordance with the price adjustment provisions specified in the SCC.

10. Payment

- 10.1. Unless otherwise specified in the SCC, payments shall be made only upon a certification by the Head of the Procuring Entity to the effect that the Goods have been rendered or delivered in accordance with the terms of this Contract and have been duly inspected and accepted. Except with the prior approval of the President no payment shall be made for services not yet rendered or for supplies and materials not yet delivered under this Contract. Ten percent (10%) of the amount of each payment shall be retained by the Procuring Entity to cover the Supplier's warranty obligations under this Contract as described in GCC Clause 17.
- 10.2. The Supplier's request(s) for payment shall be made to the Procuring Entity in writing, accompanied by an invoice describing, as appropriate, the Goods delivered and/or Services performed, and by documents submitted pursuant to the SCC provision for GCC Clause 6.2, and upon fulfillment of other obligations stipulated in this Contract.
- 10.3. Pursuant to GCC Clause 10.2, payments shall be made promptly by the Procuring Entity, but in no case later than sixty (60) days after submission of an invoice or claim by the Supplier.
- 10.4. Unless otherwise specified in the SCC, the currency in which payment is made to the Supplier under this Contract shall be in Philippine Pesos.

11. Advance Payment

- 11.1. Advance payment shall be made only after prior approval of the President, and shall not exceed fifteen percent (15%) of the Contract amount, unless otherwise directed by the President or in cases allowed under Annex "D" of RA 9184.

For Goods supplied from abroad, ten percent (10%) of the Contract price shall be paid within sixty (60) calendar days from signing of the contract and upon submission of a claim and a bank guarantee issued by a licensed bank for the equivalent amount valid until the Goods are delivered and in the form provided in All progress payments shall first be charged against the advance payment until the latter has been fully exhausted.

12. Taxes and Duties

The Supplier, whether local or foreign, shall be entirely responsible for all the necessary taxes, stamp duties, license fees, and other such levies imposed for the completion of this Contract.

13. Performance Security

- 13.1. Unless otherwise specified in the SCC, within ten (10) calendar days from receipt of the Notice of Award from the Procuring Entity but in no case later than the signing of the contract by both parties, the successful Bidder shall furnish the performance security in any the forms prescribed in the ITB Clause 33.2.

- 13.2. The performance security posted in favor of the Procuring Entity shall be forfeited in the event it is established that the winning bidder is in default in any of its obligations under the contract.
- 13.3. The performance security shall remain valid until issuance by the Procuring Entity of the Certificate of Final Acceptance.
- 13.4. Unless otherwise specified in the SCC, the performance security may be released by the Procuring Entity and returned to the Supplier after the issuance of the Certificate of Final Acceptance subject to the following conditions:
- (a) There are no pending claims against the Supplier or the surety company filed by the Procuring Entity;
 - (b) The Supplier has no pending claims for labor and materials filed against it; and
 - (c) Other terms specified in the SCC.
- 13.5. In case of a reduction of the contract value, the Procuring Entity shall allow a proportional reduction in the original performance security, provided that any such reduction is more than ten percent (10%) and that the aggregate of such reductions is not more than fifty percent (50%) of the original performance security.

14. Use of Contract Documents and Information

- 14.1. The Supplier shall not, except for purposes of performing the obligations in this Contract, without the Procuring Entity's prior written consent, disclose this Contract, or any provision thereof, or any specification, plan, drawing, pattern, sample, or information furnished by or on behalf of the Procuring Entity. Any such disclosure shall be made in confidence and shall extend only as far as may be necessary for purposes of such performance.
- 14.2. Any document, other than this Contract itself, enumerated in GCC Clause 14.1 shall remain the property of the Procuring Entity and shall be returned (all copies) to the Procuring Entity on completion of the Supplier's performance under this Contract if so required by the Procuring Entity.

15. Standards

The Goods provided under this Contract shall conform to the standards mentioned in the

Section VII. Technical Specifications; and, when no applicable standard is mentioned, to the authoritative standards appropriate to the Goods' country of origin. Such standards shall be the latest issued by the institution concerned.

16. Inspection and Tests

- 16.1. The Procuring Entity or its representative shall have the right to inspect and/or to test the Goods to confirm their conformity to the Contract specifications at no extra cost to the Procuring Entity. The SCC and
- 16.2.
- 16.3.
- 16.4. Section VII. 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.
- 16.5. If applicable, the inspections and tests may be conducted on the premises of the Supplier or its subcontractor(s), at point of delivery, and/or at the goods' final destination. If conducted on the premises of the Supplier or its subcontractor(s), all reasonable facilities and assistance, including access to drawings and production data, shall be furnished to the inspectors at no charge to the Procuring Entity.
- 16.6. The Procuring Entity or its designated representative shall be entitled to attend the tests and/or inspections referred to in this Clause provided that the Procuring Entity shall bear all of its own costs and expenses incurred in connection with such attendance including, but not limited to, all traveling and board and lodging expenses.
- 16.7. The Procuring Entity may reject any Goods or any part thereof that fail to pass any test and/or inspection or do not conform to the specifications. The Supplier shall either rectify or replace such rejected Goods or parts thereof or make alterations necessary to meet the specifications at no cost to the Procuring Entity, and shall repeat the test and/or inspection, at no cost to the Procuring Entity, upon giving a notice pursuant to GCC Clause 5.
- 16.8. The Supplier agrees that neither the execution of a test and/or inspection of the Goods or any part thereof, nor the attendance by the Procuring Entity or its representative, shall release the Supplier from any warranties or other obligations under this Contract.

17. Warranty

- 17.1. The Supplier warrants that the Goods supplied under the Contract are new, unused, of the most recent or current models, and that they incorporate all recent improvements in design and materials, except when the technical specifications required by the Procuring Entity provides otherwise.

- 17.2. The Supplier further warrants that all Goods supplied under this Contract shall have no defect, arising from design, materials, or workmanship or from any act or omission of the Supplier that may develop under normal use of the supplied Goods in the conditions prevailing in the country of final destination.
- 17.3. In order to assure that manufacturing defects shall be corrected by the Supplier, a warranty shall be required from the Supplier for a minimum period specified in the SCC. The obligation for the warranty shall be covered by, at the Supplier's option, either retention money in an amount equivalent to at least ten percent (10%) of every progress payment, or a special bank guarantee equivalent to at least ten percent (10%) of the Contract Price or other such amount if so specified in the SCC. The said amounts shall only be released after the lapse of the warranty period specified in the SCC; provided, however, that the Supplies delivered are free from patent and latent defects and all the conditions imposed under this Contract have been fully met.
- 17.4. 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, within the period specified in the SCC and with all reasonable speed, repair or replace the defective Goods or parts thereof, without cost to the Procuring Entity.
- 17.5. If the Supplier, having been notified, fails to remedy the defect(s) within the period specified in GCC Clause 17.4, the Procuring Entity may proceed to take such remedial action as may be necessary, at the Supplier's risk and expense and without prejudice to any other rights which the Procuring Entity may have against the Supplier under the Contract and under the applicable law.

18. Delays in the Supplier's Performance

- 18.1. Delivery of the Goods and/or performance of Services shall be made by the Supplier in accordance with the time schedule prescribed by the Procuring Entity in Section VI. Schedule of Requirements.
- 18.2. If at any time during the performance of this Contract, the Supplier or its Subcontractor(s) should encounter conditions impeding timely delivery of the Goods and/or performance of Services, the Supplier shall promptly notify the Procuring Entity in writing of the fact of the delay, its likely duration and its cause(s). As soon as practicable after receipt of the Supplier's notice, and upon causes provided for under GCC Clause 22, the Procuring Entity shall evaluate the situation and may extend the Supplier's time for performance, in which case the extension shall be ratified by the parties by amendment of Contract.
- 18.3. Except as provided under GCC Clause 22, a delay by the Supplier in the performance of its obligations shall render the Supplier liable to the imposition of liquidated damages pursuant to GCC Clause 19, unless an

extension of time is agreed upon pursuant to GCC Clause 29 without the application of liquidated damages.

19. Liquidated Damages

Subject to GCC Clauses 18 and 22, if the Supplier fails to satisfactorily deliver any or all of the Goods and/or to perform the Services within the period(s) specified in this Contract inclusive of duly granted time extensions if any, the Procuring Entity shall, without prejudice to its other remedies under this Contract and under the applicable law, deduct from the Contract Price, as liquidated damages, a sum equivalent to the percentage specified in the SCC of the delivered price of the delayed Goods or unperformed Services for each week or part thereof of delay until actual delivery or performance, up to a maximum deduction of the percentage specified in the SCC. Once the maximum is reached, the Procuring Entity shall rescind the Contract pursuant to GCC Clause 23, without prejudice to other courses of action and remedies open to it.

20. Settlement of Disputes

- 20.1. If any dispute or difference of any kind whatsoever shall arise between the Procuring Entity and the Supplier in connection with or arising out of this Contract, the parties shall make every effort to resolve amicably such dispute or difference by mutual consultation.
- 20.2. If after thirty (30) days, the parties have failed to resolve their dispute or difference by such mutual consultation, then either the Procuring Entity or the Supplier may give notice to the other party of its intention to commence arbitration, as hereinafter provided, as to the matter in dispute, and no arbitration in respect of this matter may be commenced unless such notice is given.
- 20.3. Any dispute or difference in respect of which a notice of intention to commence arbitration has been given in accordance with this Clause shall be settled by arbitration. Arbitration may be commenced prior to or after delivery of the Goods under this Contract.
- 20.4. Arbitration proceedings shall be conducted in accordance with the rules of procedure specified in the SCC.
- 20.5. Notwithstanding any reference to arbitration herein, the parties shall continue to perform their respective obligations under the Contract unless they otherwise agree; and the Procuring Entity shall pay the Supplier any monies due the Supplier.

21. Liability of the Supplier

- 21.1. Subject to additional provisions, if any, set forth in the SCC, the Supplier's liability under this Contract shall be as provided by the laws of the Republic of the Philippines.

- 21.2. Except in cases of criminal negligence or willful misconduct, and in the case of infringement of patent rights, if applicable, the aggregate liability of the Supplier to the Procuring Entity shall not exceed the total Contract Price, provided that this limitation shall not apply to the cost of repairing or replacing defective equipment.

22. Force Majeure

- 22.1. The Supplier shall not be liable for forfeiture of its performance security, liquidated damages, or termination for default if and to the extent that its delay in performance or other failure to perform its obligations under the Contract is the result of a *force majeure*.
- 22.2. For purposes of this Contract the terms “*force majeure*” and “fortuitous event” may be used interchangeably. In this regard, a fortuitous event or *force majeure* shall be interpreted to mean an event which the Contractor could not have foreseen, or which though foreseen, was inevitable. It shall not include ordinary unfavorable weather conditions; and any other cause the effects of which could have been avoided with the exercise of reasonable diligence by the Contractor.
- 22.3. If a *force majeure* situation arises, the Supplier shall promptly notify the Procuring Entity in writing of such condition and the cause thereof. Unless otherwise directed by the Procuring Entity in writing, the Supplier shall continue to perform its obligations under the Contract as far as is reasonably practical, and shall seek all reasonable alternative means for performance not prevented by the *force majeure*.

23. Termination for Default

- 23.1. The Procuring Entity shall terminate this Contract for default when any of the following conditions attends its implementation:
- (a) Outside of *force majeure*, the Supplier fails to deliver or perform any or all of the Goods within the period(s) specified in the contract, or within any extension thereof granted by the Procuring Entity pursuant to a request made by the Supplier prior to the delay, and such failure amounts to at least ten percent (10%) of the contract price;
 - (b) As a result of *force majeure*, the Supplier is unable to deliver or perform any or all of the Goods, amounting to at least ten percent (10%) of the contract price, for a period of not less than sixty (60) calendar days after receipt of the notice from the Procuring Entity stating that the circumstance of force majeure is deemed to have ceased; or
 - (c) The Supplier fails to perform any other obligation under the Contract.

- 23.2. In the event the Procuring Entity terminates this Contract in whole or in part, for any of the reasons provided under GCC Clauses 23 to 26, the Procuring Entity may procure, upon such terms and in such manner as it deems appropriate, Goods or Services similar to those undelivered, and the Supplier shall be liable to the Procuring Entity for any excess costs for such similar Goods or Services. However, the Supplier shall continue performance of this Contract to the extent not terminated.
- 23.3. In case the delay in the delivery of the Goods and/or performance of the Services exceeds a time duration equivalent to ten percent (10%) of the specified contract time plus any time extension duly granted to the Supplier, the Procuring Entity may terminate this Contract, forfeit the Supplier's performance security and award the same to a qualified Supplier.

24. Termination for Insolvency

The Procuring Entity shall terminate this Contract if the Supplier is declared bankrupt or insolvent as determined with finality by a court of competent jurisdiction. In this event, termination will be without compensation to the Supplier, provided that such termination will not prejudice or affect any right of action or remedy which has accrued or will accrue thereafter to the Procuring Entity and/or the Supplier.

25. Termination for Convenience

- 25.1. The Procuring Entity may terminate this Contract, in whole or in part, at any time for its convenience. The Head of the Procuring Entity may terminate a contract for the convenience of the Government if he has determined the existence of conditions that make Project Implementation economically, financially or technically impractical and/or unnecessary, such as, but not limited to, fortuitous event(s) or changes in law and national government policies.
- 25.2. The Goods that have been delivered and/or performed or are ready for delivery or performance within thirty (30) calendar days after the Supplier's receipt of Notice to Terminate shall be accepted by the Procuring Entity at the contract terms and prices. For Goods not yet performed and/or ready for delivery, the Procuring Entity may elect:
- (a) to have any portion delivered and/or performed and paid at the contract terms and prices; and/or
 - (b) to cancel the remainder and pay to the Supplier an agreed amount for partially completed and/or performed goods and for materials and parts previously procured by the Supplier.
- 25.3. If the Supplier suffers loss in its initial performance of the terminated contract, such as purchase of raw materials for goods specially manufactured for the Procuring Entity which cannot be sold in open market, it shall be allowed to recover partially from this Contract, on a *quantum meruit* basis. Before recovery may be made, the fact of loss must

be established under oath by the Supplier to the satisfaction of the Procuring Entity before recovery may be made.

26. Termination for Unlawful Acts

26.1. The Procuring Entity may terminate this Contract in case it is determined *prima facie* that the Supplier has engaged, before or during the implementation of this Contract, in unlawful deeds and behaviors relative to contract acquisition and implementation. Unlawful acts include, but are not limited to, the following:

- (a) Corrupt, fraudulent, and coercive practices as defined in ITB Clause 3.1(a);
- (b) Drawing up or using forged documents;
- (c) Using adulterated materials, means or methods, or engaging in production contrary to rules of science or the trade; and
- (d) Any other act analogous to the foregoing.

27. Procedures for Termination of Contracts

27.1. The following provisions shall govern the procedures for termination of this Contract:

- (a) Upon receipt of a written report of acts or causes which may constitute ground(s) for termination as aforementioned, or upon its own initiative, the Implementing Unit shall, within a period of seven (7) calendar days, verify the existence of such ground(s) and cause the execution of a Verified Report, with all relevant evidence attached;
- (b) Upon recommendation by the Implementing Unit, the Head of the Procuring Entity shall terminate this Contract only by a written notice to the Supplier conveying the termination of this Contract. The notice shall state:
 - (i) that this Contract is being terminated for any of the ground(s) afore-mentioned, and a statement of the acts that constitute the ground(s) constituting the same;
 - (ii) the extent of termination, whether in whole or in part;
 - (iii) an instruction to the Supplier to show cause as to why this Contract should not be terminated; and
 - (iv) special instructions of the Procuring Entity, if any.
- (c) The Notice to Terminate shall be accompanied by a copy of the Verified Report;

- (d) **Within a period of seven (7) calendar days from receipt of the Notice of Termination, the Supplier shall submit to the Head of the Procuring Entity a verified position paper stating why this Contract should not be terminated. If the Supplier fails to show cause after the lapse of the seven (7) day period, either by inaction or by default, the Head of the Procuring Entity shall issue an order terminating this Contract;**
- (e) **The Procuring Entity may, at anytime before receipt of the Supplier's verified position paper to withdraw the Notice to Terminate if it is determined that certain items or works subject of the notice had been completed, delivered, or performed before the Supplier's receipt of the notice;**
- (f) **Within a non-extendible period of ten (10) calendar days from receipt of the verified position paper, the Head of the Procuring Entity shall decide whether or not to terminate this Contract. It shall serve a written notice to the Supplier of its decision and, unless otherwise provided, this Contract is deemed terminated from receipt of the Supplier of the notice of decision. The termination shall only be based on the ground(s) stated in the Notice to Terminate;**
- (g) **The Head of the Procuring Entity may create a Contract Termination Review Committee (CTRC) to assist him in the discharge of this function. All decisions recommended by the CTRC shall be subject to the approval of the Head of the Procuring Entity; and**
- (h) **The Supplier must serve a written notice to the Procuring Entity of its intention to terminate the contract at least thirty (30) calendar days before its intended termination. The Contract is deemed terminated if it is not resumed in thirty (30) calendar days after the receipt of such notice by the Procuring Entity.**

28. Assignment of Rights

The Supplier shall not assign his rights or obligations under this Contract, in whole or in part, except with the Procuring Entity's prior written consent.

29. Contract Amendment

Subject to applicable laws, no variation in or modification of the terms of this Contract shall be made except by written amendment signed by the parties.

30. Application

These General Conditions shall apply to the extent that they are not superseded by provisions of other parts of this Contract.

Section V. Special Conditions of Contract

Special Conditions of Contract

GCC Clause	
1.1(g)	The Procuring Entity is <i>Batangas Medical Center</i>
1.1(i)	The Supplier is <i>[to be inserted at the time of contract award]</i> .
1.1(j)	<p>The Funding Source is:</p> <p>The Government of the Philippines (GOP) through <i>General Appropriations Acts of 2014</i> in the amount of <i>SEVEN MILLION FIVE HUNDRED PESOS ONLY</i></p>
1.1(k)	The Project Site is <i>Batangas Medical Center</i>
5.1	<p>The Procuring Entity's address for Notices is:</p> <p>BATANGAS MEDICAL CENTER,</p> <p>Kumintang Ilaya, Batangas City</p> <p>(043) 723-6176</p>
10.1	No further instructions
10.4	No further instructions.

13.1	No further instructions.
13.4	No further instructions.
13.4(c)	“No further instructions”.
17.3	<p><i>Select one, delete the other.</i></p> <p>Three (3) months after acceptance by the Procuring Entity of the delivered Goods or after the Goods are consumed, whichever is earlier.</p>
17.4 and 17.5	The period for correction of defects in the warranty period is <i>[insert number of days]</i> .
0	<p>The applicable rate is one tenth (1/10) of one (1) percent of the cost of the unperformed portion for every day of delay.</p> <p>The maximum deduction shall be ten percent (10%) of the amount of contract. Once the cumulative amount of liquidated damages reaches ten percent (10%) of the amount of the contract, the procuring entity shall rescind the contract, without prejudice to other courses of action and remedies open to it.</p>
20.4	In the case of a dispute between the Procuring Entity and the Supplier, the dispute shall be resolved in accordance with Republic Act 9285 (“R.A. 9285”), otherwise known as the “Alternative Dispute Resolution Act of 2004.”
21.1	“No additional provision

Section VI. Schedule of Requirements

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

[illegible]

Section VII. Technical Specifications

Technical Specifications

Item No. 1	D5% LRS	Qty.	50,000 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1 LITER BOTTLE, HARD PLASTIC			

Item No. 2	D10% Water	Qty.	1,000 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 CC BOTTLE, HARD PLASTIC			

Item No. 3	D5 0.3%, NACL	Qty.	6,000 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1 LITER BOTTLE, PLASTIC			

Item No. 4	D5% 0.3% NACL	Qty.	12,000 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 CC BOTTLE, HARD PLASTIC			

Technical Specifications

Item No. 5	D5% Ionosol MB	Qty.	300 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1 LITER BOTTLE, HARD PLASTIC			

Item No. 6	D 5% Ionosol MB	Qty.	4,000 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 CC BOTTLE, HARD PLASTIC			

Item No. 7	D 5% LRS	Qty.	4,000 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 CC BOTTLE, HARD PLASTIC			

Item No. 8	D5% Normosol M	Qty.	1,000 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1L BOTTLE, HARD PLASTIC			

Technical Specifications

Item No. 9	D5% NSS	Qty.	8,000 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1L BOTTLE, HARD PLASTIC			

Item No. 10	D5% NSS	Qty.	2,000 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 cc BOTTLE, HARD PLASTIC			

Item No. 11	D5% WATER	Qty.	6,000 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1 LITER BOTTLE, HARD PLASTIC			

Item No. 12	D5% WATER	Qty.	3,000 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250 cc BOTTLE, GLASS			

Technical Specifications

Item No. 13	D5% Water	Qty.	10,000 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 cc BOTTLE			

Item No. 14	Plain LRS	Qty.	10,000
Name of Manufacturer		Country Origin	
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1LITER BOTTLE			

Item No. 15	Plain LRS	Qty.	6,000 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 cc BOTTLE			

Item No.1 6	Plain NSS	Qty.	50,000 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1LITTER, BOTTLE			

Technical Specifications

Item No. 17	Plain NSS	Qty.	6,000 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 cc BOTTLE			

Item No. 18	Plain NSS for Irrigation	Qty.	2,000 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1 LITER BOTTLE			

Item No. 19	Acetylcysteine	Qty.	1,000 sachet
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
200 MG SACHET			

Item No. 20	Acetylcysteine	Qty.	10 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
200 MG/ML VIAL			

Technical Specifications

Item No. 21	Aciclovir	Qty.	500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
200 MG TABLET			

Item No. 22	Aciclovir	Qty.	10 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
200MG/5ML SUSPENSION,60 ML			

Item No. 23	Aciclovir	Qty.	100 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
400 MG TABLET			

Item No. 24	All in One Admixture	Qty.	80
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1,500 KILO CALORIES			

Technical Specifications

Item No. 25	Allopurinol	Qty.	4,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG TABLET			

Item No. 26	Allopurinol	Qty.	250 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
300 MG TABLET			

Item No. 27	Amikacin Sulfate	Qty.	4,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG AMPULE(IM,IV),2 ML			

Item No. 28	Amikacin Sulfate	Qty.	4,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250 MG AMPULE(IM,IV),2 ML			

Technical Specifications

Item No. 29	Amikacin Sulfate	Qty.	3,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG AMPULE(IM,IV),2 ML			

Item No. 30	Amino Acids, Crystalline	Qty.	600 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
6%, 100ML BOTTLE			

Item No. 31	Aminophylline	Qty.	100 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
25MG/ML AMPULE			

Item No. 32	Amiodarone	Qty.	500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
200 MG TABLET			

Technical Specifications

Item No. 33	Amiodarone	Qty.	50 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50MG/ML AMPULE			

Item No. 34	Amlodipine	Qty.	20,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG (AS BESILATE) TABLET			

Item No. 35	Amlodipine	Qty.	20,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
AMLODIPINE 5 MG (AS BESILATE) TABLET			

Item No. 36	Amoxicillin	Qty.	100 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100MG/5ML DROPS, 10 ML			

Technical Specifications

Item No. 37	Amoxicillin	Qty.	100 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
125 MG/ML SUSPENSION,60 ML			

Item No. 38	Amoxicillin	Qty.	100 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250MG/ML SUSPENSION,60 ML			

Item No. 39	Amoxicillin	Qty.	6,000 capsules
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG CAPSULE			

Item No. 40	Ampicillin	Qty.	1,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1 GM VIAL			

Technical Specifications

Item No. 41	Ampicillin	Qty.	20,000 capsules
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250 MG VIAL			

Item No. 42	Ampicillin	Qty.	4,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG VIAL			

Item No. 43	Ampicillin- Sulbactam	Qty.	300 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1.5GM VIAL			

Item No. 44	Ampicillin- Sulbactam	Qty.	200 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
375 GM VIAL			

Technical Specifications

Item No. 45	Ampicillin- Sulbactam	Qty.	2,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
750 MG VIAL			

Item No. 46	Anti Rabies Vaccine Serum (Equine)	Qty.	100 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
200IU/ML,5ML VIAL			

Item No. 47	Anti Tetanus 1	Qty.	10,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 AMPULE			

Item No. 48	Ascorbic Acid	Qty.	20 botlltes
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG/5ML SYRUP,60 ML			

Technical Specifications

Item No. 49	Ascorbic Acid	Qty.	20 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG/5ML DROPS,15 ML			

Item No. 50	Ascorbic Acid	Qty.	2,500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG TABLET			

Item No.51	Aspirin	Qty.	5,000 capsules
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
80 MG TABLET			

Item No. 52	Atenolol	Qty.	100 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG TABLET			

Technical Specifications

Item No. 53	Atenolol	Qty.	2,500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50 MG TABLET			

Item No. 54	Atracurium	Qty.	2,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG/ML AMPULE, 2.5 ML			

Item No. 55	Atropine Sulfate	Qty.	250 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1MG/ML AMPULE (IM/IV)			

Item No. 56	Azithromycin	Qty.	50 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
200 MG/5 ML SUSPENSION, 30ML			

Technical Specifications

Item No. 57	Azithromycin	Qty.	6,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG TABLET			

Item No. 58	Azithromycin	Qty.	300 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG VIAL			

Item No. 59	Barium Sulfate Powder, USP Grade	Qty.	500
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
340 GM			

Item No. 60	Benzathine Benzyl Penicillin	Qty.	500 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1.2 UNIT VIAL			

Technical Specifications

Item No. 61	Benzyl Penicillin	Qty.	20,000 vials
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1 MILLION UNITS VIAL			

Item No. 62	Betahistine	Qty.	500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
24 MG TABLET			

Item No. 63	Betahistine	Qty.	500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
16 MG TABLET			

Item No. 64	Biperiden	Qty.	100 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
2 MG TABLET			

Technical Specifications

Item No. 65	Biphasic Isophane Human Insulin	Qty.	250
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
70/30			

Item No. 66	Bisacodyl	Qty.	500
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
BISACODYL 10 MG SUPPOSITORY			

Item No.67	Bisacodyl	Qty.	500
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
5 MG SUPPOSITORY			

Item No. 68	Bisacodyl	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
5 MG TABLET			

Technical Specifications

Item No. 69	Bleomycin	Qty.	20 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
15MG VIAL(IM,IV,SC)			

Item No. 70	Budesonide	Qty.	2,000 nebule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250MCG/ML NEBULE			

Item No. 71	Budesonide+Formeterol	Qty.	50
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
160MCG TURBOHALER			

Item No. 72	Bupivacaine	Qty.	2,000
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
0.5% 4ML(SPINAL)W8% DEXTROSE			

Technical Specifications

Item No. 73	Bupivacaine	Qty.	1,500 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
0.5%(ISOBARIC),10ML AMPULE/VIAL			

Item No. 74	Butamirate Citrate	Qty.	2,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50 MG SR TABLET			

Item No. 75	Buthorpanol	Qty.	1,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
2MG/ML, 1ML			

Item No. 76	Calcium Carbonate	Qty.	10,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG TABLET			

Technical Specifications

Item No. 77	Calcium Carbonate	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
600 MG TABLET			

Item No. 78	Calcium Gluconate	Qty.	2,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10%, 10ML VIAL			

Item No. 79	Catopril	Qty.	4,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
25 MG TABLET			

Item No. 80	Carbamazepine	Qty.	50 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100MG/5ML SYRUP, 120ML			

Technical Specifications

Item No. 81	Carbamazepine	Qty.	2,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
200 MG TABLET			

Item No. 82	Carboplatin	Qty.	30 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
150 MG VIAL			

Item No. 83	Carboplatin	Qty.	30 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
450 MG VIAL			

Item No. 84	Castor Oil	Qty.	20
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
(USP GRADE)			

Technical Specifications

Item No.85	Cefadroxil	Qty.	500 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG CAPSULE			

Item No. 86	Cefalexin	Qty.	50 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG/5ML DROPS,15 ML DROPS			

Item No. 87	Cefalexin	Qty.	30 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
125MG/5ML SUSPENSION,60ML			

Item No. 88	Cefalexin	Qty.	30 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250MG/5ML SUSPENSION,60ML			

Technical Specifications

Item No. 89	Cefalexin	Qty.	2,500 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG CAPSULE			

Item No. 90	Cefazolin	Qty.	500 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1 GM VIAL			

Item No. 91	Cefepime	Qty.	1,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1 GM VIAL			

Item No. 92	Cefepime	Qty.	500 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG VIAL			

Technical Specifications

Item No. 93	Cefepime	Qty.	50 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG/5ML SUSPENSION,60ML			

Item No. 94	Cefepime	Qty.	3,000 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
200 MG CAPSULE			

Item No. 95	Cefepime	Qty.	20 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
20MG/5ML DROPS,10 ML			

Item No. 96	Cefotaxime	Qty.	2,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1 GM VIAL			

Technical Specifications

Item No. 97	Cefotaxime	Qty.	250 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250 MG VIAL			

Item No. 98	Cefotaxime	Qty.	500 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG VIAL			

Item No. 99	Cefoxitin	Qty.	5,000
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1 GM VIAL			

Item No. 100	Ceftazidime	Qty.	1,500 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1 GM VIAL			

Technical Specifications

Item No. 101	Ceftazidime	Qty.	1,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG VIAL			

Item No. 102	Ceftriaxone	Qty.	10,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1 GM VIAL			

Item No. 103	Ceftriaxone	Qty.	100 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250 MG VIAL			

Technical Specifications

Item No. 104	Ceftriaxone	Qty.	250 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG VIAL			

Technical Specifications

Item No. 105	Cefuroxime	Qty.	50 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
125MG/5ML SUSPENSION,60ML			

Item No. 106	Cefuroxime	Qty.	500 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250 MG VIAL			

Item No. 107	Cefuroxime	Qty.	50 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250MG/ML SUSPENSION, 60 ML			

Item No. 108	Cefuroxime	Qty.	16,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MGTABLET			

Technical Specifications

Item No. 109	Cefuroxime	Qty.	15,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
750 MG VIAL			

Item No. 110	Celecoxib	Qty.	250 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG CAPSULE			

Item No. 111	Celecoxib	Qty.	4,000 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
200 MG CAPSULE			

Item No. 112	Celecoxib	Qty.	10,000 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
400 MG CAPSULE			

Technical Specifications

Item No. 113	Citirizine	Qty.	500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG TABLET			

Item No. 114	Citirizine	Qty.	20 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
5MG/5ML SYRUP,30 ML			

Item No. 115	Chloramphenicol	Qty.	1,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1 GM VIAL			

Item No. 116	Chloramphenicol	Qty.	10 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
125MG/ML SUSPENSION,60ML			

Technical Specifications

Item No. 117	Chloramphenicol	Qty.	500 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG CAPSULE			

Item No. 118	Cinnarizine	Qty.	500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
25 MG TABLET			

Item No. 119	Cinnarizine	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
75 MG			

Item No. 120	Ciprofloxacin	Qty.	6,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
200 MG/ML VIAL			

Technical Specifications

Item No. 121	Ciprofloxacin	Qty.	6,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG TABLET			

Item No. 122	Cisplatin	Qty.	30 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50 MG VIAL			

Item No. 123	Clarithromycin	Qty.	50 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
125 MG/ML SUSPENSION, 50 ML			

Item No. 124	Methyldopa	Qty.	2,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
125MG TABLET			

Technical Specifications

Item No. 125	Clarithromycin	Qty.	5,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG TABLET			

Item No. 126	Clindamycin	Qty.	2,000 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
150 MG CAPSULE			

Item No. 127	Clindamycin	Qty.	5,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
150MG/ML AMPULE,2 ML			

Technical Specifications

Item No. 128	Clindamycin	Qty.	15,000 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
300 MG CAPSULE			

Item No. 129	Clindamycin	Qty.	10 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
75MG/ML SUSPENSION,60ML			

Item No. 130	Cotrimoxazole	Qty.	10 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
200MG+40MG/5ML SUSP,60ML			

Item No. 131	Clonidine	Qty.	8,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
75 MG TABLET			

Technical Specifications

Item No. 132	Clopidogrel	Qty.	8,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
75 MG TABLET			

Item No. 133	Cloxacillin	Qty.	50 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
125 MG/ML SUSPENSION, 60 ML			

Item No. 134	Cloxacillin	Qty.	1,000 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250 MG CAPSULE			

Item No. 135	Cloxacillin	Qty.	50 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250MG/5ML SUSPENSION,60ML			

Technical Specifications

Item No. 136	Cloxacillin	Qty.	15,000 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG CAPSULE			

Item No. 137	Co-Amoxiclav	Qty.	20 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
228MG+62.5 MG/5ML SUSPENSION,70ML			

Item No. 138	Co-Amoxiclav	Qty.	20 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
CO-AMOXICLAV 312MG/5ML SUSPENSION,60ML			

Item No. 139	Co-Amoxiclav	Qty.	6,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
625 MG TABLET			

Technical Specifications

Item No. 140	Colchicine	Qty.	4,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG TABLET			

Item No. 141	Cotrimoxazole	Qty.	50 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
400 MG TABLET			

Item No. 142	Cotrimoxazole	Qty.	10 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
400MG+80MG/5ML SUSP,60ML			

Item No. 143	Cotrimoxazole	Qty.	50 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
800 MG TABLET			

Technical Specifications

Item No. 144	Cyclophosphamide	Qty.	500 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
200 MG VIAL			

Item No. 145	Cyclophosphamide	Qty.	500 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG VIAL			

Item No. 146	Cytarabine	Qty.	100 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100MG/ML VIAL			

Item No. 147	Dexamethazone	Qty.	500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
4 MG TABLET			

Technical Specifications

Item No. 148	Dexamethazone	Qty.	1,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
4MG/ML AMPULE,1 ML(IM,IV)			

Item No. 149	Dextromethorpan	Qty.	50 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG TABLET			

Item No. 150	Dextromethorpan	Qty.	10 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
5MG/5ML SYRUP,60 ML			

Item No. 151	Dextrose	Qty.	2,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50 % 50 cc VIAL			

Technical Specifications

Item No. 152	Diazepam	Qty.	250 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG TABLET			

Item No. 153	Diazepam	Qty.	250 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
5 MG TABLET			

Item No. 154	Diazepam	Qty.	4,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
5MG/ML, 2ML AMPULE(IM,IV)			

Item No. 155	Diclofenac	Qty.	50 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
25MG/ML AMPULE(IM/IV/SC)			

Technical Specifications

Item No. 156	Diclofenac	Qty.	100 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
75 MG TABLET			

Item No. 157	Digoxin	Qty.	2,500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250 MCG TABLET			

Item No. 158	Digoxin	Qty.	2,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250MCG/ML AMPULE			

Item No. 159	Diltiazem	Qty.	250 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
HCL 60 MG TABLET			

Technical Specifications

Item No. 160	Diltiazem	Qty.	250 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
HCL 30 MG TABLET			

Item No. 161	Diphenhydramine	Qty.	10 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
12.5MG/5ML SYRUP,60ML			

Item No. 162	Diphenhydramine	Qty.	1,000 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50 MG CAPSULE			

Item No. 163	Diphenhydramine	Qty.	1,500 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50 MG/ML AMPULE(IM/IV)			

Technical Specifications

Item No. 164	Dobutamine	Qty.	600 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
2MG/ML,250MLD5W PREMIXED BOTTLE			

Item No. 165	Dobutamine	Qty.	1,500 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50 MG/ML AMPULE/VIAL			

Item No. 166	Docetaxel	Qty.	20 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
20MG/0.5ML VIAL			

Item No. 167	Docetaxel	Qty.	20 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
40MG/ML,2ML VIAL			

Technical Specifications

Item No. 168	Domperidone	Qty.	500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG TABLET			

Item No. 169	Domperidone	Qty.	20 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1MG/ML, 30 ML SUSPENSION			

Item No. 170	Dopamine	Qty.	250 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
40MG/ML,5ML AMPULE			

Item No. 171	Dopamine	Qty.	750 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
800MCG/ML, 250ML PREMIXED BOTTLE			

Technical Specifications

Item No. 172	Doxorubicin	Qty.	1,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50 MG VIAL			

Item No. 173	Doxorubicin	Qty.	1,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG VIAL			

Item No. 174	Doxycycline	Qty.	3,000 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG CAPSULE			

Item No. 175	Enalapril	Qty.	500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG TABLET			

Technical Specifications

Item No. 176	Enalapril	Qty.	500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
5 MG TABLET			

Item No. 177	Enoxaparin	Qty.	2,500 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100MG/ML, 0.4 cc AMPULE, 100MG/ML 0.4CC			

Item No. 178	Ephedrine Sulfate	Qty.	1,200 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50MG/ML AMPULE(IM/IV)			

Item No. 179	Ephedrine	Qty.	6,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1 MG/ML AMPULE			

Technical Specifications

Item No. 180	Epoetin Alfa (Recombinant Human Erythropoetin)	Qty.	6,000
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
4000 IU			

Item No. 181	Ertapenem	Qty.	10 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
(AS SODIUM) 1 GM VIAL			

Item No. 182	Erythromycin	Qty.	50 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
200MG/5ML SUSPENSION,60ML			

Item No. 183	Erythromycin	Qty.	50 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
40MG/5ML SUSPENSION,30ML			

Technical Specifications

Item No. 184	Erythromycin	Qty.	500 tubes
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
OINTMENT 0.5% ,5 GM TUBE			

Item No. 185	Etoposide	Qty.	30 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG VIAL			

Item No. 186	Famotidine	Qty.	500 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
20 MG LYOPHILIZED POWDER VIAL			

Item No. 187	Famotidine	Qty.	250 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
20 MG TABLET			

Technical Specifications

Item No. 188	Felodipine	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG MR TABLET			

Item No. 189	Felodipine	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
5 MG MR TABLET			

Item No. 190	Fenofibrate	Qty.	250 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
160 MG MR TABLET			

Item No. 191	Fenofibrate	Qty.	250 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
200 MG TABLET			

Technical Specifications

Item No. 192	Fentanyl Citrate	Qty.	2,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50MCG/ML AMPULE,2 ML			

Item No. 193	Ferrous sulfate	Qty.	100 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
150MG/5ML,60 ML SYRUP			

Item No. 194	Ferrous sulfate	Qty.	100 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
15MG/0.6ML,15 ML DROPS			

Item No. 195	Ferrous sulfate+Folic Acid	Qty.	15,000
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
60 MG ELEMETAL IRON			

Technical Specifications

Item No. 196	Fluconazole	Qty.	500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
150 MG TABLET			

Item No. 197	Fluconazole	Qty.	250 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50 MG TABLET			

Item No. 198	Flunarizine HCL	Qty.	250 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
5 MG CAPSULE			

Item No. 199	Fluorouracil	Qty.	250 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50MG/ML VIAL			

Technical Specifications

Item No. 200	Fluticasone+Salmeterol	Qty.	100
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250MCG+25MCG 120 DOSES			

Item No. 201	Fluticasone+Salmeterol	Qty.	200
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250MCG+50MCG 120 DOSES			

Item No. 202	Folic Acid	Qty.	4,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
5 MG TABLET			

Item No. 203	Furosemide	Qty.	20,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG/ML AMPULE, 2ML			

Technical Specifications

Item No. 204	Furosemide	Qty.	6,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
FUROSEMIDE 20 MG TABLET			

Item No. 205	Furosemide	Qty.	5,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
40 MG TABLET			

Item No. 206	Fusidate Sodium Fusidic Acid	Qty.	100 tube
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
2%,5 G TUBE			

Item No. 207	Gabapentin	Qty.	100 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG CAPSULE			

Technical Specifications

Item No. 208	Gabapentin	Qty.	100 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
300 MG CAPSULE			

Item No. 209	Sodium Chloride	Qty.	5,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
0.9% , 50 cc VIAL			

Item No. 210	Gentamicin	Qty.	10,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
40 MG/ML VIAL,2 ML			

Item No. 211	Glicazide	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
60MG/MR TABLET			

Technical Specifications

Item No. 212	Glicazide	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
80 MG TABLET			

Item No. 213	Glipizide	Qty.	500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
5 MG TABLET			

Item No. 214	Glycerin	Qty.	100
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
2 GM SUPPOSITORY(RECTAL)			

Item No. 215	Glycerin Trinitrate	Qty.	100
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
5 MG PATCH			

Technical Specifications

Item No. 216	Haloperidol	Qty.	10 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
5 MG/ML AMPULE, 1 ML			

Item No. 217	Heparin	Qty.	3,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1,000 UNIT VIAL			

Item No. 218	Hepatitis B Vaccine	Qty.	20
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10MCG/0.5ML MONODOSE(IM)			

Item No. 219	Human Albumin	Qty.	250 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
20 % BOTTLE, 100 ML			

Technical Specifications

Item No. 220	Hydralazine	Qty.	3,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
20 MG/ML AMPULE			

Item No. 221	Hydralazine	Qty.	250 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
25 MG TABLET			

Item No. 222	Hydralazine	Qty.	250 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50 MG TABLET			

Item No. 223	Hydrocortizone	Qty.	10,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG VIAL			

Technical Specifications

Item No. 224	Hydrocortizone	Qty.	2,500 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250 MG VIAL			

Item No. 225	Hydrogen Peroxide	Qty.	2,500 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
3%, 120 ML BOTTLE			

Item No. 226	HydroxynEthyl Starch	Qty.	500 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
6 % BOTTLE			

Item No. 227	Hydroxyzine	Qty.	50 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50 MG AMPULE			

Technical Specifications

Item No. 228	Hyoscine	Qty.	2,500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG TABLET			

Item No. 229	Hyoscine	Qty.	5,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
20 MG/ML AMPULE			

Item No. 230	Ibuprofen	Qty.	100 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG/5ML SYRUP BOTTLE,60ML			

Item No. 231	Ifosfamide	Qty.	30 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
2 GM VIAL			

Technical Specifications

Item No. 232	Immunoglobulin Normal, Human	Qty.	10 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50 MG VIAL			

Item No. 233	Influenza Polyvalent Vaccine	Qty.	50 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
0.5 vial+Pre-filled Syringe Diluent (IM)			

Item No. 234	ING75MG+Rifampicin150Mg +PZA400+Ethambutol	Qty.	250 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
275MG TABLET			

Item No. 235	ING75MG+Rifampicin150Mg +Ethambutol	Qty.	200 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
275 MG TABLET			

Technical Specifications

Item No. 236	Insulin, Regular	Qty.	250 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100IU,10 ML VIAL			

Item No. 237	Intraocular Irrigating Solution	Qty.	150
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 ML(BALANCED SALT SOLUTION)			

Item No. 238	Ipratropium+Salbutamol	Qty.	20,000 nebule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MCG+2.5MG/2ML NEBULE			

Item No. 239	Isoniazid	Qty.	10 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
200MG/ML SYRUP			

Technical Specifications

Item No. 240	Isophane Human Insulin	Qty.	1,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 IU/ML VIAL			

Item No. 241	Isosorbide 5 Mononitrate	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
20 MG TABLET			

Item No. 242	Isosorbide 5 Mononitrate	Qty.	500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
40 MG TABLET			

Item No. 243	Isosorbide Dinitrate	Qty.	1,000 tablet
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG TABLET			

Technical Specifications

Item No. 244	Isosorbide Dinitrate	Qty.	250 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1MG/ML AMPULE			

Item No. 245	Isosorbide Dinitrate	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
20 MG TABLET			

Item No. 246	Isosorbide Dinitrate	Qty.	1,000 tablet
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
5 MG TABLET			

Item No. 247	Isoxsuprine HCL	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG TABLET			

Technical Specifications

Item No. 248	Isoxsuprine HCL	Qty.	1,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
5MG/ML AMPULE			

Item No. 249	Ketamine	Qty.	200 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50 MG/ML VIAL(IM/IV)			

Item No. 250	Ketoconazole	Qty.	200 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
200 MG TABLET			

Item No. 251	Ketorolac	Qty.	6,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG TABLET			

Technical Specifications

Item No. 252	Ketorolac	Qty.	20,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
30MG/ML AMPULE (IM/IV)			

Item No. 253	Lactulose	Qty.	1,500 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
3.3MG/5ML (66%) 120 ML BOTTLE			

Item No. 254	Lagundi	Qty.	10 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
300MG/5ML SYRUP, 60 ML BOTTLE			

Item No. 255	Lagundi	Qty.	150 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
600 MG TABLET			

Technical Specifications

Item No. 256	L-Aspariginase	Qty.	50 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10,000 U VIAL			

Item No. 257	Levofloxacin	Qty.	4,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG TABLET			

Item No. 258	Levofloxacin	Qty.	1,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
5MG/ML VIAL			

Item No. 259	Levofloxacin	Qty.	2,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
750 MG TABLET			

Technical Specifications

Item No. 260	Levothyroxine	Qty.	100 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG TABLET			

Item No. 261	Levothyroxine	Qty.	100 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
25 MCG TABLET			

Item No. 262	Levothyroxine	Qty.	100 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50 MCG TABLET			

Item No. 263	Lidocaine	Qty.	40 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10% 50 ML SPRAY			

Technical Specifications

Item No. 264	Lidocaine	Qty.	2,000
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
2%,1.8ML(W/ EPINEPHRINE)CARPULE			

Item No. 265	Lidocaine HCL	Qty.	6,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
2%, 5ML POLYAMPULE			

Item No. 266	Lidocaine Jelly	Qty.	2 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
2% 30GMS			

Item No. 267	Loperamide	Qty.	500 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
2 MG CAPSULE			

Technical Specifications

Item No. 268	Loratadine	Qty.	250 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG TABLET			

Item No. 269	Loratadine	Qty.	10 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
5MG/5ML,30 ML SYRUP			

Item No. 270	Losartan	Qty.	5,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG TABLET			

Item No. 271	Losartan	Qty.	10,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG W/HCTZ TABLET			

Technical Specifications

Item No. 272	Losartan	Qty.	4,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50 MG TABLET			

Item No. 273	Losartan	Qty.	5,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50MG W/HCTZ TABLET			

Item No. 274	Magnesium Sulfate	Qty.	2,500 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250MG/ML AMPULE/VIAL(M/IV),10ML			

Item No. 275	Mannitol	Qty.	5,000 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
20% BOTTLE, 500 ML BOTTLE			

Technical Specifications

Item No. 276	Mebendazole	Qty.	20 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100MG/5ML SUSPENSION,30ML BOTTLE			

Item No. 277	Mebendazole	Qty.	50 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
MEBENDAZOLE 500 MG TABLET			

Item No. 278	Mefinamic Acid	Qty.	1,000 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250 MG CAPSULE			

Item No. 279	Mefinamic Acid	Qty.	20 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50 MG/5ML SUSPENSION,60ML BOTTLE			

Technical Specifications

Item No. 280	Mefinamic Acid	Qty.	4,000 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG CAPSULE			

Item No. 281	Mercaptopurine	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50 MG TABLET			

Item No. 282	Meropenem	Qty.	4,200 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1 GM VIAL			

Item No. 283	Meropenem	Qty.	4,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG VIAL			

Technical Specifications

Item No. 284	Mesna	Qty.	20
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG/ML,5 ML			

Item No. 285	Metformin HCL	Qty.	5,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG TABLET			

Item No. 286	Methothrexate	Qty.	500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
2.5 MG TABLET			

Item No. 287	Methotrexate	Qty.	50 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
METHOTREXATE 25MG/5ML,2 ML VIAL			

Technical Specifications

Item No. 288	Methyldopa	Qty.	2,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250 MG TABLET			

Item No. 289	Methylergometrine	Qty.	10,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
125 MCG TABLET			

Item No. 290	Methylergometrine	Qty.	4,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
200MCG/ML AMPULE			

Item No. 291	Methylprednisolone	Qty.	4 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1 GM/16ML+DILUENT VIAL			

Technical Specifications

Item No. 292	Metoclopramide	Qty.	500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG TABLET			

Item No. 293	Metoclopramide	Qty.	10 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
5MG/5ML SYRUP,60 ML BOTTLE			

Item No. 294	Metoclopramide	Qty.	2,500 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
5MG/ML, 2 ML AMPULE			

Item No. 295	Metoprolol	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG TABLET			

Technical Specifications

Item No. 296	Metoprolol	Qty.	5,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50 MG TABLET			

Item No. 297	Metronidazole	Qty.	250
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1 GM SUPPOSITORY			

Technical Specifications

Item No. 298	Metronidazole	Qty.	20 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
125 MG/ML SUSPENSION,60 ML			

Item No. 299	Metronidazole	Qty.	4,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG TABLET			

Technical Specifications

Item No. 300	Metronidazole	Qty.	20,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
5MG/ML, 100 ML VIAL			

Item No. 301	Midazolam	Qty.	50 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
15 MG TABLET			

Item No. 302	Midazolam	Qty.	2,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
5MG/ML AMPULE(IM/IV),1ML			

Technical Specifications

Item No. 303	Modified Fluid Gelatin	Qty.	500 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
4% BOTTLE			

Technical Specifications

Item No. 304	Monobasic/Dibasic Sodium Phospate	Qty.	500 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
19G/7G/133ML			

Item No. 305	Monobasic/Dibasic Sodium Phospate	Qty.	500 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
48G/18G/100ML SOLN,45ML BOTTLE			

Item No. 306	Montelukast	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG TABLET			

Item No. 307	Montelukast	Qty.	250 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
4 MG TABLET,CHEWABLE			

Technical Specifications

Item No. 308	Montelukast	Qty.	250 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
5 MG TABLET,CHEWABLE			

Item No. 309	Morphine Sulfate	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG TABLET			

Item No. 310	Morphine Sulfate	Qty.	1,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10MG/ML AMPULE(IM/IV/SC),1ML			

Technical Specifications

Item No. 311	Morphine Sulfate	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
30 MG TABLET			

Technical Specifications

Item No. 312	Multi Vitamins	Qty.	5,000 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
CAPSULE			

Item No. 313	Multi Vitamins	Qty.	100 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
DROPS			

Item No. 314	Multi Vitamins	Qty.	100 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
SYRUP			

Item No. 315	Mupirocin	Qty.	1,000 tube
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
2%/ 5 GM OINTMENT TUBE			

Technical Specifications

Item No. 316	Nalbuphine	Qty.	3,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG/ML, 1ML AMPULE(IM/IV/SC)			

Item No. 317	Naloxone HCL	Qty.	10 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
400MCG/ML,1ML AMPULE			

Item No. 318	Naproxen Sodium	Qty.	500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
550 MG TABLET			

Item No. 319	Neostigmine	Qty.	10 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MCG/ML AMPULE(IM/IV/SC)			

Technical Specifications

Item No. 320	Nicardipine	Qty.	50 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1 MG/ML AMPULE, 2ML			

Item No. 321	Nicardipine	Qty.	1,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1MG/ML AMPULE, 10ML			

Item No. 322	Nifedipine	Qty.	250 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG TABLET			

Item No. 323	Nifedipine	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
30 MG/MR TABLET			

Technical Specifications

Item No. 324	Nifedipine	Qty.	250 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
5 MG TABLET			

Item No. 325	Nimodipine	Qty.	250 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
30 MG TABLET			

Item No. 326	Nitrofurantoin	Qty.	250 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG CAPSULE			

Item No. 327	IO Pomidol	Qty.	800 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
NON-IONIC CONTRAST MEDIA, VIAL, G12MG/ML EQUIV., to 300MG IODINE/ML 50ML			

Technical Specifications

Item No. 328	Norpinephrine	Qty.	2,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
AMPULE, 1MG/ML (BITARTRATE)			

Item No. 329	Norfloxacin	Qty.	100 tablet
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
200 MG TABLET			

Item No. 330	Nystatin	Qty.	4 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100,000 U/ML SUSPENSION,30ML			

Item No. 331	Ofloxacin	Qty.	500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
200 MG TABLET			

Technical Specifications

Item No. 332	Omeprazole	Qty.	2,500 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
20 MG CAPSULE			

Item No. 333	Omeprazole	Qty.	4,500 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
40 MG CAPSULE			

Item No. 334	Omeprazole	Qty.	12,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
40 MG POWDER+10ML SOLVENT AMPULE			

Item No. 335	Ondansetron	Qty.	500 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
2MG/ML,4ML AMPULE			

Technical Specifications

Item No. 336	Oral Rehydration Salts	Qty.	500 sachet
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
SACHET			

Item No. 337	Oxantel+Pyrantel	Qty.	50 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG TABLET			

Item No. 338	Oxantel+Pyrantel	Qty.	10
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG(EMBONATE)5ML,10ML			

Item No. 339	Oxycodone	Qty.	50 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG TABLET			

Technical Specifications

Item No. 340	Oxycodone	Qty.	50 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
20 MG TABLET			

Technical Specifications

Item No. 341	Oxytocin	Qty.	5,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 IU AMPULE			

Item No. 342	Paclitaxel	Qty.	20 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
6MG/ML,5ML VIAL			

Item No. 343	Pancuronium	Qty.	4 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
2MG/ML AMPULE,2ML			

Technical Specifications

Item No. 344	Paracetamol	Qty.	305 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100MG/ML DROPS,15ML(ALCOHOL FREE)			

Item No. 345	Paracetamol	Qty.	100 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10MG/ML,100 ML VIAL			

Item No. 346	Paracetamol	Qty.	150
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
125 MG SUPPOSITORY			

Item No. 347	Paracetamol	Qty.	600 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
125MG/ML SYRUP,60ML(125MG/ML)(ALCOHOL FREE)			

Technical Specifications

Item No. 348	Paracetamol	Qty.	16,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
150MG/ML,AMPULE			

Item No. 349	Paracetamol	Qty.	500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250 MG TABLET			

Item No. 350	Paracetamol	Qty.	10,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG TABLET			

Item No. 351	Penicillin Benzathine	Qty.	250 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1.2 MU			

Technical Specifications

Item No. 352	Penicillin Crystalline	Qty.	15,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1 MU VIAL			

Item No. 353	Pethidine	Qty.	10 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50G/ML AMPULE(IM/IV/SC),1ML			

Item No. 354	Phenobarbital	Qty.	800 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
130MG/ML AMPULE(IM/IV),1ML			

Item No. 355	Phenobarbital	Qty.	500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
15 MG TABLET			

Technical Specifications

Item No. 356	Phenobarbital	Qty.	500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
30 MG TABLET			

Item No. 357	Phenobarbital	Qty.	500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
60 MG TABLET			

Item No. 358	Phenytoin	Qty.	15,000 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG CAPSULE			

Item No. 359	Phenytoin	Qty.	10 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
30MG/5ML SUSPENSION, 120 ML BOTTLE			

Technical Specifications

Item No. 360	Phenytoin	Qty.	3,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50MG/ML AMPULE,2ML(IM/IV),2ML			

Item No. 361	Phenytoin	Qty.	500 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50MG/ML,5 ML AMPULE			

Item No. 362	Phytomenadione	Qty.	5,000
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG/ML AMPULE(IM/IV/SC)			

Item No. 363	Piperacillin Tazobactam	Qty.	4,750 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
2.25 GM VIAL			

Technical Specifications

Item No. 364	Piperacillin Tazobactam	Qty.	5,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
4.5 GM VIAL			

Item No. 365	Pneumococcal	Qty.	20 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
VACCINE			

Item No. 366	Potassium Chloride	Qty.	6,000 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
2 MEQ/ML VIAL			

Item No. 367	Potassium Chloride	Qty.	4,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
750 MG DURULES TABLET			

Technical Specifications

Item No. 368	Prednisone	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG TABLET			

Item No. 369	Prednisone	Qty.	250 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
20 MG TABLET			

Technical Specifications

Item No.370	Prednisone	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
5 MG TABLET			

Item No. 371	Propanolol	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG TABLET			

Technical Specifications

Item No. 372	Propanolol	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
40 MG TABLET			

Item No. 373	Propofol	Qty.	1,500 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10MG/ML, 20ML AMPULE/VIAL			

Item No. 374	Propofol	Qty.	20
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10MG/ML, 50ML PREFILLED SYRINGE			

Item No.375	Propylthiouracil	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50 MG TABLET			

Technical Specifications

Item No. 376	Pyrazinamide	Qty.	20 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250MG/ML SYRUP,60 ML BOTTLE			

Item No. 377	Rabies Vaccine Chick Embryo Cell	Qty.	60 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
2.5IU/ML, 1 DOSE VIAL			

Item No.378	Ranitidine	Qty.	250 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
150 MG TABLET			

Item No. 379	Ranitidine	Qty.	4,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
25 MG/ML AMPULE			

Technical Specifications

Item No. 380	Rifampicin	Qty.	10 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
200 MG/5ML SYRUP			

Item No. 381	Rocuronium	Qty.	20 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10MG/ML VIAL,2.5ML VIAL			

Item No. 382	Ropivacaine	Qty.	6 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10MG/ML,10 ML AMPULE			

Item No. 383	Rosuvastatin	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
10 MG TABLET (CALCIUM)			

Technical Specifications

Item No. 384	Rosuvastatin	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
20 MG TABLET (CALCIUM)			

Item No. 385	Salbutamol	Qty.	15 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
2MG/5ML, 60 ML SYRUP			

Item No. 386	Salbutamol	Qty.	1,000 nebules
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
2MG/ML, 2.5 ML NEBULE (AS SULFATE)			

Item No. 387	Sambong	Qty.	250 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG TABLET			

Technical Specifications

Item No. 388	Silver Sulfadiazine cream	Qty.	400
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
,25 GMS			

Item No. 389	Simvastatin	Qty.	4,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
20 MG TABLET			

Item No. 390	Simvastatin	Qty.	5,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
40 MG TABLET			

Item No. 391	Sodium Bicarbonate	Qty.	300 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1 MEQ/ML VIAL,50 ML			

Technical Specifications

Item No. 392	Sodium Chloride	Qty.	500 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
2.5 MEQ/ML VIAL, 20ML			

Item No. 393	Somatostatin	Qty.	4 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250 MCG AMPULE			

Item No. 394	Spironolactone	Qty.	250 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG TABLET			

Item No. 395	Spironolactone	Qty.	500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
25 MG TABLET			

Technical Specifications

Item No. 396	Spironolactone	Qty.	500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50 MG TABLET			

Item No. 397	Sterile Water of Injection	Qty.	14,000 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 ML BOTTLE			

Item No. 398	Streptokinase	Qty.	10 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1,500,000 VIAL			

Item No. 399	Streptomycin Sulfate	Qty.	50 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1 GM VIAL			

Technical Specifications

Item No. 400	Suxametonium	Qty.	500 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
20MG/ML,10 ML VIAL			

Item No. 401	Tamoxifen	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
TAMOXIFEN 20 MG TABLET			

Item No. 402	Telmisartan	Qty.	100 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
40 MG TABLET			

Technical Specifications

Item No. 403	Telmisartan	Qty.	100 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
80 MG TABLET			

Technical Specifications

Item No. 404	Terbutaline S04	Qty.	10 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MCG/ML AMPULE(IM/IV/SC),1ML			

Item No. 405	Tetanus Toxoid	Qty.	3,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
AMPULE			

Item No. 406	Tetracycline	Qty.	50 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250 MG CAPSULE			

Item No. 407	Tetracycline	Qty.	50 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG CAPSULE			

Technical Specifications

Item No. 408	Theophylline	Qty.	6 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
25MG/5ML(26.7MG/5ML)60ML SYRUP			

Item No. 409	Tobramycin	Qty.	150 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
0.3%,5ML BOTTLE EYE DROPS			

Item No. 410	Tobramycin	Qty.	150 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
0.3%+DEXAMETHASONE0.1%,5ML EYE DROPS,BOTTLE			

Item No.411	Tramadol	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG MR TABLET			

Technical Specifications

Item No. 412	Tramadol	Qty.	3,600 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50 MG TABLET			

Item No. 413	Tramadol	Qty.	14,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
50MG/ML AMPULE,1 ML(IM,IV,SC)			

Item No. 414	Tranexamic Acid	Qty.	7,000 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
100 MG/ML AMPULE,5ML			

Item No. 415	Tranexamic Acid	Qty.	5,000 capsule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG CAPSULE			

Technical Specifications

Item No. 416	Trimetazidine	Qty.	1,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
20 MG TABLET			

Item No.417	Trimetazidine	Qty.	1,200 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
35 MG TABLET			

Item No. 418	Valproate Sodium	Qty.	500 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250 MG TABLET			

Technical Specifications

Item No. 419	Valproic Acid	Qty.	500 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
250MG/5ML, 120 ML SYRUP			

Technical Specifications

Item No. 420	Vancomycin	Qty.	600 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
500 MG VIAL			

Item No. 421	Verapamil	Qty.	10 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
2.5MG/ML,2ML AMPULE			

Item No. 422	Verapamil	Qty.	50 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
40 MG TABLET			

Technical Specification

Item No. 423	Verapamil	Qty.	50 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
80 MG TABLET			

Item No. 424	Vincristine	Qty.	20 vial
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
1MG/ML,2ML VIAL			

Item No.425	Vitamin B Complex	Qty.	15,000 tablets
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
TABLET			

Technical Specification

Item No. 426	Vitamin B1 B6 B12	Qty.	400 ampule
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
AMPULE			

Item No. 427	Zinc	Qty.	150 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
27.5MG/ML, 15ML DROPS BOTTLE			

Item No. 428	Zinc	Qty.	200 bottles
Name of Manufacturer		Country Origin	
Brand:			
PURCHASER'S SPECIFICATION		SUPPLIER'S SPECIFICATION	
55MG/5ML, 60ML SYRUP BOTTLE			

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Section VIII. Bidding Forms

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Bid Form

Date: _____
Invitation to Bid² N°: _____

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

Gentlemen and/or Ladies:

Having examined the Bidding Documents including 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 Bidding Documents for the sum of *[total Bid amount in words and figures]* or such other sums as may be ascertained in accordance with the Schedule of Prices attached herewith and made part of this Bid.

We undertake, if our Bid is accepted, to deliver the goods in accordance with the delivery schedule specified in the Schedule of Requirements.

If our Bid is accepted, we undertake to provide a performance security in the form, amounts, and within the times specified in the Bidding Documents.

We agree to abide by this Bid for the Bid Validity Period specified in **BDS** provision for **ITB** Clause 18.2 and it shall remain binding upon us and may be accepted at any time before the expiration of that period.

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:³

Name and address of agent	Amount and Currency	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 or any Bid you may receive.

We certify/confirm that we comply with the eligibility requirements as per **ITB** Clause 5 of the Bidding Documents.

² If ADB, JBIC and WB funded projects, use IFB.

³ Applicable only if the Funding Source is the ADB, JBIC or WB.

Dated this _____ day of _____ 20_____.

[signature]

[in the capacity of]

Duly authorized to sign Bid for and on behalf of _____

For Goods Offered From Abroad

Name of Bidder _____. Invitation to Bid⁴ Number _____. 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)

⁴ If ADB, JBIC and WB funded projects, use IFB.

For Goods Offered From Within the Philippines

Name of Bidder _____. Invitation to Bid⁵ Number __. Page . of ____.

1	2	3	4	5	6	7	8	9	10
Item	Description	Country of origin	Quantity	Unit price EXW per item	Cost of local labor, raw material, and component ²	Total price EXW per item (cols. 4 x 5)	Unit prices per item final destination and unit price of other incidental services	Sales and other taxes payable per item if Contract is awarded	Total Price delivered Final Destination (col 8 + 9) x 4

⁵ If ADB, JBIC and WB funded projects, use IFB.

Contract Agreement Form

THIS AGREEMENT made the ____ day of _____ 20____ between *[name of PROCURING ENTITY]* of the Philippines (hereinafter called “the Entity”) of the one part and *[name of Supplier]* of *[city and country of Supplier]* (hereinafter called “the Supplier”) of the other part:

WHEREAS the Entity invited Bids for certain goods and ancillary services, viz., *[brief description of goods and services]* and has accepted a Bid by the Supplier for the supply of those goods and services in the sum of *[contract price in words and figures]* (hereinafter called “the Contract Price”).

NOW THIS AGREEMENT WITNESSETH AS FOLLOWS:

1. In this Agreement words and expressions shall have the same meanings as are respectively assigned to them in the Conditions of Contract referred to.
2. The following documents shall be deemed to form and be read and construed as part of this Agreement, viz.:
 - (a) the Bid Form and the Price Schedule submitted by the Bidder;
 - (b) the Schedule of Requirements;
 - (c) the Technical Specifications;
 - (d) the General Conditions of Contract;
 - (e) the Special Conditions of Contract; and
 - (f) the Entity’s Notification of Award.
3. In consideration of the payments to be made by the Entity to the Supplier as hereinafter mentioned, the Supplier hereby covenants with the Entity to provide the goods and services and to remedy defects therein in conformity in all respects with the provisions of the Contract
4. The Entity hereby covenants to pay the Supplier in consideration of the provision of the goods and services and the remedying of defects therein, the Contract Price or such other sum as may become payable under the provisions of the contract at the time and in the manner prescribed by the contract.

IN WITNESS whereof the parties hereto have caused this Agreement to be executed in accordance with the laws of the Republic of the Philippines on the day and year first above written.

Signed, sealed, delivered by _____ the _____ (for the Entity)

Error! No bookmark name given. Signed, sealed, delivered by _____ the _____ (for the Supplier).

Omnibus Sworn Statement

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 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 of *[Name of Bidder]*, I have full power and authority to do, execute and perform any and all acts necessary to represent it in the bidding for *[Name of the Project]* of the *[Name of the Procuring Entity]*;

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 and/or to represent the *[Name of Bidder]* in the bidding as shown in the attached *[state title of attached document showing proof of authorization (e.g., duly notarized Secretary's Certificate issued by the corporation or the members of the joint venture)]*;

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;
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: I am 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 following responsibilities as a Bidder:
 - a) Carefully examine all of the Bidding Documents;
 - b) Acknowledge all conditions, local or otherwise, affecting the implementation of the Contract;
 - c) Made an estimate of the facilities available and needed for the contract to be bid, if any; and
 - d) Inquire or secure Supplemental/Bid Bulletin(s) issued for the *[Name of the Project]*.

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

Bidder's Representative/Authorized Signatory

[JURAT]

Bank Guarantee Form for Advance Payment

To: *[name and address of PROCURING ENTITY]*
[name of Contract]

Gentlemen and/or Ladies:

In accordance with the payment provision included in the Special Conditions of Contract, which amends Clause 10 of the General Conditions of Contract to provide for advance payment, *[name and address of Supplier]* (hereinafter called the "Supplier") shall deposit with the PROCURING ENTITY a bank guarantee to guarantee its proper and faithful performance under the said Clause of the Contract in an amount of *[amount of guarantee in figures and words]*.

We, the *[bank or financial institution]*, as instructed by the Supplier, agree unconditionally and irrevocably to guarantee as primary obligator and not as surety merely, the payment to the PROCURING ENTITY on its first demand without whatsoever right of objection on our part and without its first claim to the Supplier, in the amount not exceeding *[amount of guarantee in figures and words]*.

We further agree that no change or addition to or other modification of the terms of the Contract to be performed thereunder or of any of the Contract documents which may be made between the PROCURING ENTITY and the Supplier, shall in any way release us from any liability under this guarantee, and we hereby waive notice of any such change, addition, or modification.

This guarantee shall remain valid and in full effect from the date of the advance payment received by the Supplier under the Contract until *[date]*.

Yours truly,

Signature and seal of the Guarantors

[name of bank or financial institution]

[address]

[date]

REPUBLIC OF THE PHILIPPINES)
CITY OF _____) S.S.
X-----X

BID-SECURING DECLARATION

Invitation to Bid/Request for Expression of Interest No.1: *[Insert reference number]*

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

I/We², the undersigned, declare that:

1. I/We understand that, according to your conditions, bids must be supported by a Bid Security, which may be in the form of a Bid-Securing Declaration.
2. I/We accept that: (a) I/we will be automatically disqualified from bidding for any contract with any procuring entity for a period of two (2) years upon receipt of your Blacklisting Order; and, (b) I/we will pay the applicable fine provided under Section 6 of the Guidelines on the Use of Bid Securing Declaration³, if I/we have committed any of the following actions:
 - (i) Withdrawn my/our Bid during the period of bid validity required in the Bidding Documents; or
 - (ii) Fail or refuse to accept the award and enter into contract or perform any and all acts necessary to the execution of the Contract, in accordance with the Bidding Documents after having been notified of your acceptance of our Bid during the period of bid validity.
3. I/We understand that this Bid-Securing Declaration shall cease to be valid on the following circumstances:
 - (a) Upon expiration of the bid validity period, or any extension thereof pursuant to your request;
 - (b) I am/we are declared ineligible or post-disqualified upon receipt of your notice to such effect, and (i) I/we failed to timely file a request for reconsideration or (ii) I/we filed a waiver to avail of said right;
 - (c) I am/we are declared as the bidder with the Lowest Calculated and Responsive Bid/Highest Rated and Responsive Bid⁴, and I/we have furnished the performance security and signed the Contract.

GPPB Resolution No. 03-2012,

