BIDDING DOCUMENTS

for

(EPA 2025) SUPPLY AND DELIVERY OF VETERINARY DRUGS AND SUPPLIES FOR PCC-OED

under

PB 2024-78

Preface

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

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

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

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

a. All the documents listed in the Table of Contents are normally required for the procurement of Goods. However, they should be adapted as necessary to the circumstances of the particular Procurement Project.

b. Specific details, such as the "*name of the Procuring Entity*" and "*address for bid submission*," should be furnished in the Instructions to Bidders, Bid Data Sheet, and Special Conditions of Contract. The final documents should contain neither blank spaces nor options.

c. This Preface and the footnotes or notes in italics included in the Invitation to Bid, Bid Data Sheet, General Conditions of Contract, Special Conditions of Contract, Schedule of Requirements, and Specifications are not part of the text of the final document, although they contain instructions that the Procuring Entity should strictly follow.

d. The cover should be modified as required to identify the Bidding Documents as to the Procurement Project, Project Identification Number, and Procuring Entity, in addition to the date of issue.

e. Modifications for specific Procurement Project details should be provided in the Special Conditions of Contract as amendments to the Conditions of Contract. For easy completion, whenever reference has to be made to specific clauses in the Bid Data Sheet or Special Conditions of Contract, these terms shall be printed in bold typeface on Sections I (Instructions to Bidders) and III (General Conditions of Contract), respectively.

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

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Glossary of Acronyms, Terms, and Abbreviations

ABC – Approved Budget for the Contract.

BAC – Bids and Awards Committee.

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

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

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

BIR – Bureau of Internal Revenue.

BSP – Bangko Sentral ng Pilipinas.

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

CDA - Cooperative Development Authority.

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

CIF – Cost Insurance and Freight.

CIP – Carriage and Insurance Paid.

CPI – Consumer Price Index.

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

DTI – Department of Trade and Industry.

EXW – Ex works.

FCA – "Free Carrier" shipping point.

FOB – "Free on Board" shipping point.

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

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

GFI – Government Financial Institution.

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

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

GOP – Government of the Philippines.

GPPB – Government Procurement Policy Board.

INCOTERMS – International Commercial Terms.

Infrastructure Projects – Include the construction, improvement, rehabilitation, demolition, repair, restoration or maintenance of roads and bridges, railways, airports, seaports, communication facilities, civil works components of information technology projects, irrigation, flood control and drainage, water supply, sanitation, sewerage and solid waste management systems, shore protection, energy/power and electrification facilities, national buildings, school buildings, hospital buildings, and other related construction projects of the government. Also referred to as *civil works or works*. (2016 revised IRR, Section 5[u])

LGUs – Local Government Units.

NFCC – Net Financial Contracting Capacity.

NGA – National Government Agency.

PhilGEPS - Philippine Government Electronic Procurement System.

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

PSA – Philippine Statistics Authority.

SEC – Securities and Exchange Commission.

SLCC – Single Largest Completed Contract.

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

UN – United Nations.

Section I. Invitation to Bid

Notes on the Invitation to Bid

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

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

a. The date of availability of the Bidding Documents, which shall be from the time the IB is first advertised/posted until the deadline for the submission and receipt of bids;

b. The place where the Bidding Documents may be acquired or the website where it may be downloaded;

c. The deadline for the submission and receipt of bids; and

d. Any important bid evaluation criteria (*e.g.*, the application of a margin of preference in bid evaluation).

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

INVITATION TO BID
ITB/Identification No. PB 2024-78

(EPA 2025) SUPPLY AND DELIVERY OF VETERINARY DRUGS AND SUPPLIES FOR PCC-OED

1. The **PHILIPPINE CARABAO CENTER**, through the **GAA Fund** intends to apply the sum of **ELEVEN MILLION FIVE HUNDRED THIRTY-ONE THOUSAND NINE HUNDRED FOURTEEN PESOS ONLY(P 11,531,914.00)** being the ABC to payments under the contract for **(EPA 2025) SUPPLY AND DELIVERY OF VETERINARY DRUGS AND SUPPLIES FOR PCC-OED** under **ITB/Identification No. PB 2024-78** Bids received in excess of the ABC shall be automatically rejected at bid opening.

2. The **PHILIPPINE CARABAO CENTER** now invites bids for the above Procurement Project. Completion of delivery/Works/Services is required within **Sixty (60) Calendar days** as specified in the Schedule of Requirements. Bidders should have completed, within **Five (5) years** from the date of submission and receipt of bids, a contract similar to the Project. The description of an eligible bidder is contained in the Bidding Documents, particularly, in Section II (Instructions to Bidders).

3. Bidding will be conducted through open competitive bidding procedures using a non-discretionary "pass/fail" criterion as specified in the 2016 revised Implementing Rules and Regulations (IRR) of Republic Act (RA) No. 9184.

   Bidding is restricted to Filipino citizens/sole proprietorships, partnerships, or organizations with at least sixty percent (60%) interest or outstanding capital stock belonging to citizens of the Philippines, and to citizens or organizations of a country the laws or regulations of which grant similar rights or privileges to Filipino citizens, pursuant to RA No. 5183.

4. Prospective Bidders may obtain further information from **PHILIPPINE CARABAO CENTER** and inspect the Bidding Documents at the address given below during Monday to Friday (8am-5pm) except weekends and holidays.

5. A complete set of Bidding Documents may be acquired by interested Bidders on – **October 22 – November 18, 2024,** from the given address and website(s) below and upon payment of the applicable fee for the Bidding Documents, pursuant to the latest Guidelines issued by the GPPB, in the amount of **Eleven Thousand Five Hundred Pesos Only (P 11,500.00) non-refundable**. The Procuring Entity shall allow the bidder to present its proof of payment for the fees presented in person, by facsimile, or through electronic means not later than the submission of their bids.

   We encourage bidders to download the bidding documents through PhilGEPS or the PCC Website and pay through bank. Please coordinate with the BAC Secretariat Office through the email address and contact number given below for the bank detail procedures.

NATIONAL HEADQUARTERS AND GENE POOL Science City of Muñoz, Nueva Ecija, Philippines, 3120 • Tel No 044 456 0731 • URL: http://pcc.gov.ph • Email: oed@pcc.gov.ph **MANILA LIAISON OFFICE** 2nd Floor, ICC Building, National Irrigation Administration Complex, EDSA, Quezon City, Philippines, 0830 • Tel/Fax No.: (02) 8927 7270 • Email: mlo@pcc.gov.ph

6. The **PHILIPPINE CARABAO CENTER** will hold a Pre-Bid Conference (via Zoom for bidders) on **November 6, 2024 (10:00 AM)** at **PHILIPPINE CARABAO CENTER, National Headquarter and Gene Pool,** Science City of Muñoz, Nueva Ecija through web conference platform via Zoom. A Personal Meeting ID will be sent through email for each participant at least one day before the event. Prospective bidders need to signify their intention to participate by sending an email to bac-secretariat@pcc.gov.ph

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7. Bids must be duly received by the BAC Secretariat through manual submission at the office address indicated below on or before **November 18, 2024, until 5:00 PM**. Late bids shall not be accepted.

8. All Bids must be accompanied by a bid security in any of the acceptable forms and in the amount stated in **ITB** Clause 14.

9. Bid opening shall be on **November 19, 2024 (10:00 AM)** at the given address below through web conference platform via Zoom (face to face for the BAC).

   A Personal Meeting ID will be sent to participants through email who submitted bid proposals.

10. The **PHILIPPINE CARABAO CENTER** reserves the right to reject any and all bids, declare a failure of bidding, or not award the contract at any time prior to contract award in accordance with Sections 35.6 and 41 of the 2016 revised IRR of RA No. 9184, without thereby incurring any liability to the affected bidder or bidders.

11. The PHILIPPINE CARABAO CENTER Bids and Awards Committee (BAC) will use a non-discretionary and non-discriminatory measure based on sheer luck or chance, which is "draw lots", in the event that two or more bidders have been post-qualified and determined as the bidder having the Lowest Calculated Responsive Bid (LCRB) to determine the final LCRB (In alphabetical order, the bidders shall pick one rolled paper. The lucky bidder who would pick the paper with "Congratulations" remark shall be declared as the final bidder having LCRB and recommended for award of the contract.)

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12. Please refer to the following schedule of activities:

Activity	Date	Remarks
Availability of Bidding documents	**October 22 – November 18, 2024**, (Except Holidays, Saturdays and Sundays) 8:00 AM to 5:00 PM	Downloadable through PhilGEPS, PCC Website and can be requested through email (**bac-secretariat@pcc.gov.ph**)
Payment of Bidding Documents (before submission of bid proposals)	**October 22 – November 18, 2024**, (Except Holidays, Saturdays and Sundays) 8:00 AM to 5:00 PM	Prospective Bidders may pay the bidding documents through bank deposit to: Bank Name/Branch: Land Bank of the Philippines-CLSU Account Name: PHILIPPINE CARABAO CENTER LBP Branch where the deposit was made: If bidders opted to pay cash, request Statement of Account (SOA) first from PCC Accounting Office before proceeding to the PCC Cashier's Office Prospective Bidders must email the scanned copy of deposit slip to BAC Secretariat Office with the following details: ▪ Project Title ▪ Company Name with Contact Information BAC Secretariat Office: **bac-secretariat@pcc.gov.ph** PCC Cashier's Office will issue Official Receipt after the validation/confirmation of the bank deposit.
Pre-bid Conference (Online for bidders)	**November 6, 2024** (10:00 AM)	A Personal Meeting ID will be sent by email for each participant at least one day before the event. Prospective bidders need to signify their intention to participate by sending an email to **bac-secretariat@pcc.gov.ph**
Deadline for the submission of bids	**November 18, 2024** (Until 5:00 PM)	**Late bids shall not be accepted.** Through personal delivery or through courier at the PCC BAC Secretariat Office Bidders must ensure that the Bid Proposals are properly delivered and received by the BAC Secretariat Office on or before the deadline of submission.
Bid Opening (Online for bidders)	**November 19, 2024** (10:00 AM)	A Personal Meeting ID will be sent to participants through email who submitted bid proposals on time.

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13. The PHILIPPINE CARABAO CENTER (PCC) does not condone any forms of solicitation on any prospective winning and losing bidders by any of our staff/employees or any other party. Any sort of this kind shall be reported immediately to PCC (oed@pccgov.ph).

14. For further information, please refer to

MS. NOEMI V. BALAIS
Head, BAC Secretariat Office
PHILIPPINE CARABAO CENTER
National Headquarters and Gene Pool
Science City of Munoz, Nueva Ecija
Email: bac-secretariat@pcc.gov.ph
Contact No.: 0917 824 4374
Website: www.pcc.gov.ph

ERIC P. PALACPAC
Chairperson, Bids and Awards Committee
October 21, 2024

Section II. Instructions to Bidders

Notes on the Instructions to Bidders

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

1. Scope of Bid

The Procuring Entity, **PHILIPPINE CARABAO CENTER** wishes to receive Bids for the **(EPA 2025) SUPPLY AND DELIVERY OF VETERINARY DRUGS AND SUPPLIES FOR PCC-OED**, with identification number **PB 2024-78.**

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

The Procurement Project (referred to herein as "Project") is composed of **(EPA 2025) SUPPLY AND DELIVERY OF VETERINARY DRUGS AND SUPPLIES FOR PCC-OED** the details of which are described in Section VII (Technical Specifications).

2. Funding Information

2.1. The GOP through the source of funding as indicated below for *[indicate funding year]* in the amount of **P 11,531,914.00**

2.2. The source of funding is **GAA Fund**

3. Bidding Requirements

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

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

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

4. Corrupt, Fraudulent, Collusive, and Coercive Practices

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

5. Eligible Bidders

5.1. Only Bids of Bidders found to be legally, technically, and financially capable will be evaluated.

5.2. Foreign ownership exceeding those allowed under the rules may participate pursuant to:

i. When a Treaty or International or Executive Agreement as provided in Section 4 of the RA No. 9184 and its 2016 revised IRR allow foreign bidders to participate;

ii. 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;

iii. When the Goods sought to be procured are not available from local suppliers; or

iv. When there is a need to prevent situations that defeat competition or restrain trade.

a. Foreign ownership limited to those allowed under the rules may participate in this Project.

5.3. Pursuant to Section 23.4.1.3 of the 2016 revised IRR of RA No.9184, the Bidder shall have an SLCC that is at least one (1) contract similar to the Project the value of which, adjusted to current prices using the PSA's CPI, must be at least equivalent to:

a. For the procurement of Non-expendable Supplies and Services: The Bidder must have completed a single contract that is similar to this Project, equivalent to at least **twenty-five percent (25%)** of the ABC amounting to **P 2,882,978.50 or aggregate contracts (refer to Bid Data Sheet)**.

5.4. The Bidders shall comply with the eligibility criteria under Section 23.4.1 of the 2016 IRR of RA No. 9184.

6. Origin of Goods

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

7. Subcontracts

7.1. **Subcontracting is not allowed**.

7.2. Subcontracting of any portion of the Project does not relieve the Supplier of any liability or obligation under the 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.

8. Pre-Bid Conference

The Procuring Entity will hold a pre-bid conference for this Project on the specified date and time and either at its physical address **AFMD Conference Room Philippine Carabao Center, National Headquarters and Gene Pool, Science City of Muñoz, Nueva Ecija** and/or through videoconferencing/webcasting} as indicated in paragraph 6 of the **IB.**

9. Clarification and Amendment of Bidding Documents

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

10. Documents comprising the Bid: Eligibility and Technical Components

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

10.2. The Bidder's SLCC as indicated in **ITB** Clause 5.3 should have been completed within **Five (5) years** prior to the deadline for the submission and receipt of bids.

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

11. Documents comprising the Bid: Financial Component

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

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

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

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

12. Bid Prices

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

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

i. The price of the Goods quoted EXW (ex-works, ex-factory, ex-warehouse, ex-showroom, or off-the-shelf, as applicable);

ii. The cost of all customs duties and sales and other taxes already paid or payable;

iii. The cost of transportation, insurance, and other costs incidental to delivery of the Goods to their final destination; and

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

b. For Goods offered from abroad:

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

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

13. Bid and Payment Currencies

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

13.2. Payment of the contract price shall be made in:

a. **Philippine Pesos.**

14. Bid Security

14.1. The Bidder shall submit a Bid Securing Declaration[1] or any form of Bid Security in the amount indicated in the **BDS**, which shall be not less than the percentage of the ABC in accordance with the schedule in the **BDS**.

14.2. The Bid and bid security shall be valid until **One Hundred Twenty (120) calendar days from the date of Bid Opening**. Any Bid not accompanied by an acceptable bid security shall be rejected by the Procuring Entity as non-responsive.

15. Sealing and Marking of Bids

Each Bidder shall submit one copy of the first and second components of its Bid.
The Procuring Entity may request additional hard copies and/or electronic copies of the Bid. However, failure of the Bidders to comply with the said request shall not be a ground for disqualification.

If the Procuring Entity allows the submission of bids through online submission or any other electronic means, the Bidder shall submit an electronic copy of its Bid, which must be digitally signed. An electronic copy that cannot be opened or is corrupted shall be considered non-responsive and, thus, automatically disqualified.

16. Deadline for Submission of Bids

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

17. Opening and Preliminary Examination of Bids

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

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

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

18. Domestic Preference

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

19. Detailed Evaluation and Comparison of Bids

19.1. The Procuring Entity's BAC shall immediately conduct a detailed evaluation of all Bids rated "*passed*," using non-discretionary pass/fail criteria. The BAC shall consider the conditions in the evaluation of Bids under Section 32.2 of the 2016 revised IRR of RA No. 9184.

19.2. If the Project allows partial bids, bidders may submit a proposal on any of the lots or items, and evaluation will be undertaken on a per lot or item basis, as the case maybe. In this case, the Bid Security as required by **ITB** Clause 14 shall be submitted for each lot or item separately.

19.3. The descriptions of the lots or items shall be indicated in **Section VII (Technical Specifications)**, although the ABCs of these lots or items are indicated in the **BDS** for purposes of the NFCC computation pursuant to Section 23.4.2.6 of the 2016 revised IRR of RA No. 9184. The NFCC must be sufficient for the total of the ABCs for all the lots or items participated in by the prospective Bidder.

19.4. The Project shall be awarded as follows:

Option 1 – One Project having several items that shall be awarded as one contract.

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

20. Post-Qualification

20.1. Within a non-extendible period of five (5) calendar days from receipt by the Bidder of the notice from the BAC that it submitted the Lowest Calculated Bid, or in the case of multi-year Framework Agreement, that it is one of the eligible bidders who have submitted bids that are found to be technically and financially compliant,}the Bidder shall submit its latest income and business tax returns filed and paid through the BIR Electronic Filing and Payment System (eFPS) and other appropriate licenses and permits required by law and stated in the **BDS**.

21. Signing of the Contract

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

Section III. Bid Data Sheet

Notes on the Bid Data Sheet

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

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

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

a. Information that specifies and complements provisions of the ITB must be incorporated.

b. Amendments and/or supplements, if any, to provisions of the ITB as necessitated by the circumstances of the specific procurement, must also be incorporated.

Bid Data Sheet

ITB Clause	
5.3	For this purpose, contracts similar to the Project shall be: The Bidder must have a **Single Largest Completed Contract (SLCC)** within the last five (5) years equivalent to at least **25%** of the total Approved Budget for the Contract OR In view of the determination by the Procuring Entity that imposition of the provisions of Section 23.5.1.3 of the IRR of RA 9184 will likely result to "failure of bidding' or monopoly that will defeat the purpose of public bidding", the Bidder should comply with the following. Requirements: *Completed at least two (2) similar contracts the aggregate amount of which should be equivalent to at least "fifty percent (50%)" in the case of Non-expendable Supplies of the ABC for this Project However, in the case of Expendable Supplies, said SLCC must be at least twenty-five percent (25%) of the ABC. The largest of these similar contracts must be equivalent to at least half of the percentage of the ABC as required above.* **Similar Contract: PROJECT/S THAT INVOLVES SUPPLY AND DELIVERY OF VETERINARY DRUGS, BIOLOGICALS, VETERINARY SUPPLIES, AI SUPPLIES AND OTHER RELATED PARAPHERNALIA'S**
7.1	**Subcontracting is not allowed.**
12	The price of the Goods shall be quoted delivered duty paid (DDP) to the **PHILIPPINE CARABAO CENTER.**
14.1	The bid security shall be in the form of a Bid Securing Declaration, or any of the following forms and amounts: a. The amount of not less than **P 230,638.28** equivalent to **two percent (2%)** of ABC, if bid security is in cash, cashier's/manager's check, bank draft/guarantee or irrevocable letter of credit; or b. The amount of not less than **P 576,595.70** equivalent to **five percent (5%)** of ABC if bid security is in Surety Bond.
19.3	The project will be awarded **PER ITEM**
20.2	Post-Qualification documents 1. Latest income and business tax returns filed and paid through the BIR Electronic Filing and Payment System (eFPS) 2. Other appropriate licenses and permits as required by law The bidder with the Lowest Calculated and Responsive Bid (LCRB)/Single Calculated and Responsive Bid (SCRB) must submit the document/s to the BAC Secretariat Office within five (5) calendar days from the bid opening
21.2	No other contract documents required.

Section IV. General Conditions of Contract

Notes on the General Conditions of Contract

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

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

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

1. Scope of Contract

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

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

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

2. Advance Payment and Terms of Payment

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

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

3. Performance Security

Within ten (10) calendar days from receipt of the Notice of Award by the Bidder from the Procuring Entity but in no case later than the signing of the Contract by both parties, the successful Bidder shall furnish the performance security in any of the forms prescribed in Section 39 of the 2016 revised IRR of RA No. 9184.*{[Include if Framework Agreement will be used:] In the case o*f Framework Agreement, the Bidder may opt to furnish the performance security or a Performance Securing Declaration as defined under the Guidelines on the Use of Framework Agreement.*}*

4. Inspection and Tests

The Procuring Entity or its representative shall have the right to inspect and/or to test the Goods to confirm their conformity to the Project *{[Include if Framework Agreement will be used:]* or Framework Agreement*}* specifications at no extra cost to the Procuring Entity in accordance with the Generic Procurement Manual. In addition to tests in the **SCC**, **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.

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

5. Warranty

5.1 In order to assure that manufacturing defects shall be corrected by the Supplier, a warranty shall be required from the Supplier as provided under Section 62.1 of the 2016 revised IRR of RA No. 9184.

5.2 The Procuring Entity shall promptly notify the Supplier in writing of any claims arising under this warranty. Upon receipt of such notice, the Supplier shall, repair or replace the defective Goods or parts thereof without cost to the Procuring Entity, pursuant to the Generic Procurement Manual.

6. Liability of the Supplier

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

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

Section V. Special Conditions of Contract

Notes on the Special Conditions of Contract

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

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

a. Information that complements provisions of the GCC must be incorporated.

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

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

Special Conditions of Contract

GCC Clause	
	Delivery and Documents – For purposes of the Contract, "EXW," "FOB," "FCA," "CIF," "CIP," "DDP" and other trade terms used to describe the obligations of the parties shall have the meanings assigned to them by the current edition of INCOTERMS published by the International Chamber of Commerce, Paris. The Delivery terms of this Contract shall be as follows: *[For Goods supplied from abroad, state:]* "The delivery terms applicable to the Contract are DDP delivered **Philippine Carabao Center National Head Quarter and Genepool, Science City of Muñoz, Nueva Ecija.** in accordance with INCOTERMS." *[For Goods supplied from within the Philippines, state:*] "The delivery terms applicable to this Contract are delivered **Philippine Carabao Center National Head Quarter and Genepool, Science City of Muñoz, Nueva Ecija.** Risk and title will pass from the Supplier to the Procuring Entity upon receipt and final acceptance of the Goods at their final destination." Delivery of the Goods shall be made by the Supplier in accordance with the terms specified in Section VI (Schedule of Requirements). For purposes of this Clause the Procuring Entity's Representative at the Project site is **DR. LIZA G. BATTAD** or her authorized representatives. **Incidental Services –** The Supplier is required to provide all of the following services, including additional services, if any, specified in Section VI. Schedule of Requirements: *Select appropriate requirements and delete the rest.* a. performance or supervision of on-site assembly and/or start-up of the supplied Goods; b. furnishing of tools required for assembly and/or maintenance of the supplied Goods; c. furnishing of a detailed operations and maintenance manual for each appropriate unit of the supplied Goods; d. performance or supervision or maintenance and/or repair of the supplied Goods, for a period of time agreed by the parties, provided that this service shall not relieve the Supplier of any warranty obligations under this Contract; and

	e. training of the Procuring Entity's personnel, at the Supplier's plant and/or on-site, in assembly, start-up, operation, maintenance, and/or repair of the supplied Goods. f. *[Specify additional incidental service requirements, as needed.]* The Contract price for the Goods shall include the prices charged by the Supplier for incidental services and shall not exceed the prevailing rates charged to other parties by the Supplier for similar services. **Spare Parts –** The Supplier is required to provide all of the following materials, notifications, and information pertaining to spare parts manufactured or distributed by the Supplier: *Select appropriate requirements and delete the rest.* 1. such spare parts as the Procuring Entity may elect to purchase from the Supplier, provided that this election shall not relieve the Supplier of any warranty obligations under this Contract; and 2. in the event of termination of production of the spare parts: i. advance notification to the Procuring Entity of the pending termination, in sufficient time to permit the Procuring Entity to procure needed requirements; and ii. following such termination, furnishing at no cost to the Procuring Entity, the blueprints, drawings, and specifications of the spare parts, if requested. The spare parts and other components required are listed in **Section VI (Schedule of Requirements)** and the costs thereof are included in the contract price. The Supplier shall carry sufficient inventories to assure ex-stock supply of consumable spare parts or components for the Goods for a period of [*indicate here the time period specified. If not used indicate a time period of three times the warranty period*]. Spare parts or components shall be supplied as promptly as possible, but in any case, within [*insert appropriate time period*] months of placing the order.
	Packaging – The Supplier shall provide such packaging of the Goods as is required to prevent their damage or deterioration during transit to their final destination, as indicated in this Contract. The packaging shall be sufficient to withstand, without limitation, rough handling during transit and exposure to extreme temperatures, salt and

	precipitation during transit, and open storage. Packaging case size and weights shall take into consideration, where appropriate, the remoteness of the Goods' final destination and the absence of heavy handling facilities at all points in transit. The packaging, marking, and documentation within and outside the packages shall comply strictly with such special requirements as shall be expressly provided for in the Contract, including additional requirements, if any, specified below, and in any subsequent instructions ordered by the Procuring Entity. The outer packaging must be clearly marked on at least four (4) sides as follows: Name of the Procuring Entity Name of the Supplier Contract Description Final Destination Gross weight Any special lifting instructions Any special handling instructions Any relevant HAZCHEM classifications
	A packaging list identifying the contents and quantities of the package is to be placed on an accessible point of the outer packaging if practical. If not practical the packaging list is to be placed inside the outer packaging but outside the secondary packaging. **Transportation –** Where the Supplier is required under Contract to deliver the Goods CIF, CIP, or DDP, transport of the Goods to the port of destination or such other named place of destination in the Philippines, as shall be specified in this Contract, shall be arranged and paid for by the Supplier, and the cost thereof shall be included in the Contract Price. Where the Supplier is required under this Contract to transport the Goods to a specified place of destination within the Philippines, defined as the Project Site, transport to such place of destination in the Philippines, including insurance and storage, as shall be specified in this Contract, shall be arranged by the Supplier, and related costs shall be included in the contract price.

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

Section VI. Schedule of Requirements

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

Item No.	Description	Unit	Qty	Total ABC	Delivered, Weeks/Months/Years	End-user
1.	AI ALPHA SHEATH	pack	300	247,500.00	60 calendar days upon receipt of the Notice to Proceed (NTP)	CALF
2.	AI GLOVES	box	500	192,500.00		CALF
3.	AI GUN	piece	130	271,414.00		CALF
4.	AI STRAW SHEATHS	pack	400	62,480.00		CALF
5.	ARTIFICIAL VAGINA FOR CATTLE	piece	2	16,000.00		CRU
6.	ARTIFICIAL VAGINA FOR GOAT	piece	2	16,000.00		CRU
7.	BULL HOLDER WITH CHAIN	piece	2	7,800.00		ABGS
8.	BULL RING APPLICATOR	piece	2	6,600.00		ABGS
9.	CIDAR APPLICATOR	piece	5	7,500.00		RPS
10.	CRYO GRIPPER FOR CRYOTANKS	piece	2	5,000.00		CRU
11.	DRENCHING GUN	piece	2	11,726.00		NDBH
12.	FORCEP	piece	80	68,200.00		CALF
13.	GOBLET	bag	2	6,000.00		CRU
14.	MILK DIPPER	lot	8	160,000.00		ABGS
15.	ORAL CANNULA / ORAL DRENCHER	piece	30	10,500.00		NDBH

16.	STAINLESS TONG	piece	3	450.00		ABGS
17.	STRAW CUTTER	piece	120	78,000.00		CALF
18.	EAR TAG SUPPLIES	lot	1	1,034,450.00		ABGS
19.	ALBENDAZOLE 15%	bot	50	65,000.00		BES
20.	ALBENDAZOLE 15%	bot	100	130,000.00		CALF
21.	ALBENDAZOLE 15%	bot	190	247,000.00		NDBH
22.	AMINO ACID INJECTABLES / DEXTROLYTE	bot	20	12,000.00		LBRAF
23.	ATROPINE SULFATE - 15mg 50ml/ bottle - Minimum shelf life: 24 months from date of delivery	bot	4	2,552.00		NDBH
24.	CALCIUM BOROGLUCONATE (CBG)	bot	20	8,000.00		LBRAF
25.	CALIFORNIA MASTITIS TEST REAGENT	bot	20	24,000.00		BES
26.	CAUSTIC SODA FLAKES	bag	12	20,400.00		GP
27.	CEFTIOFUR HCL/ SODIUM INJECTABLE	bot	15	25,050.00		BES
28.	CEFTIOFUR HCL/ SODIUM INJECTABLE	bot	40	66,800.00		NDBH
29.	CHLORTETRACYCLINE HYDROXIDE WOUND SPRAY	can	10	7,500.00		BES
30.	CHLORTETRACYCLINE HYDROXIDE WOUND SPRAY	can	7	5,250.00		LBRAF
31.	DCM	bot	30	22,500.00		LBRAF

32.	DCM	bot	30	22,500.00		NDBH
33.	DEXAMETHASONE SODIUM PHOSPHATE	bot	20	8,000.00		BES
34.	DEXAMETHASONE SODIUM PHOSPHATE	bot	30	12,000.00		NDBH
35.	DISINFECTANT	bot	16	46,400.00		LBRAF
36.	DORAMECTIN INJECTABLE	bot	10	20,000.00		BES
37.	DOXYCYCLINE	bot	10	8,000.00		BES
38.	EPINEPHRINE HYDROCHLORIDE	box	4	836.00		NDBH
39.	FERRIC HYDROIDE + VITAMIN B 12 10%, INJECTABLE	bot	20	8,000.00		LBRAF
40.	HEMORRHAGIC SEPTECEMIA VACCINE	dose	650	42,250.00		NDBH
41.	HEPARIN SOLUTION, RX	bot	2	70,000.00		RPS
42.	INTRAMMAMMARY ANTIBIOTIC INFUSION TUBES FOR DRY ANIMALS	carton	1	7,100.00		BES
43.	INTRAMMAMMARY ANTIBIOTIC INFUSION TUBES FOR LACTATING ANIMALS	carton	1	7,100.00		BES
44.	IVERMECTIN INJECTABLE	bot	10	9,000.00		BES
45.	IVERMECTIN INJECTABLE	bot	7	5,950.00		NDBH
46.	IVERMECTIN INJECTABLE	bot	150	127,500.00		CALF
47.	KETOPROFEN	bot	15	17,280.00		LBRAF

48.	KETOPROFEN	bot	40	46,080.00		NDBH
49.	L-CARNITINE USP,	box	8	32,000.00		RPS
50.	LACTATED RINGER'S SOLUTION, RX	piece	12	3,840.00		RPS
51.	LIDOCAINE HYDROCHORIDE 2%	bot	20	5,000.00		NDBH
52.	LIVER TONIC	bot	10	17,600.00		BES
53.	LIVER TONIC	bot	10	17,600.00		LBRAF
54.	LIVER TONIC	bot	30	52,800.00		NDBH
55.	MINERAL BLOCK	box	10	40,000.00		LBRAF
56.	ORAL MULTIVITAMINS	bot	10	43,750.00		BES
57.	ORAL MULTIVITAMINS	bot	20	87,500.00		LBRAF
58.	OXYTETRACYCLINE LA 20%	bot	10	6,000.00		BES
59.	OXYTETRACYCLINE LA 20%	bot	35	14,945.00		NDBH
60.	PENTASULFATE SOLUTION	bot	10	15,000.00		BES
61.	PERMETHRIN SCREWWORM AEROSOL	can	12	17,496.00		NDBH
62.	POVIDONE IODINE (10% solution)	gallon	5	5,585.00		NDBH
63.	SEALING POWDER	bot	5	42,500.00		CALF
64.	TEAT SEALANT	pail	1	30,000.00		BES
65.	TOLTAZURIL	bot	10	60,000.00		BES

66.	VITAMIN ADE INJECTABLE	bot	100	160,000.00		BES
67.	VITAMIN ADE INJECTABLE	bot	135	216,000.00		CALF
68.	VITAMIN ADE INJECTABLE	bot	35	56,000.00		LBRAF
69.	VITAMIN ADE INJECTABLE	bot	105	168,000.00		NDBH
70.	VITAMIN B COMPLEX WITH LIVER EXTRACT	bot	60	23,460.00		NDBH
71.	VITAMIN B12 BUTAPHOSPHAN , INJECTABLE	bot	60	60,000.00		BES
72.	VITAMIN B12 BUTAPHOSPHAN , INJECTABLE	bot	30	30,000.00		LBRAF
73.	AVIAN PPD TUBERCULIN ANTIGEN	dose	200	34,000.00		BES
74.	BOVINE PPD TUBERCULIN ANTIGEN	dose	3,500	280,000.00		BES
75.	BOVINE PPD TUBERCULIN ANTIGEN	dose	470	37,600.00		NDBH
76.	BRUCELLA TEST ANTIGEN	bot	10	88,000.00		BES
77.	BRUCELLA TEST ANTIGEN	bot	2	17,600.00		NDBH
78.	CIDAR	pack	25	360,000.00		RPS
79.	FASCIOLICIDAL DRUG	bot	25	183,750.00		BES
80.	FASCIOLICIDAL DRUG	bot	20	147,000.00		LBRAF
81.	GONADOTROPIN RELEASING HORMONE	vial	22	132,000.00		RPS

82.	GONADOTROPIN RELEASING HORMONE	vial	50	300,000.00		CALF
83.	HEMORRHAGIC SEPTECEMIA VACCINE	dose	5,000	325,000.00		BES
84.	HORMONE, HUMAN CHORIONIC GONADOTROPIN (hCG)	box	15	292,500.00		CALF
85.	HORMONE, HUMAN CHORIONIC GONADOTROPIN (hCG)	box	12	234,000.00		RPS
86.	PAG BLOOD KIT	kit	1	157,000.00		RPS
87.	PAG MILK KIT	kit	1	132,000.00		RPS
88.	PORCINE PITUITARY-DERIVED FOLLICLE STIMULATING HORMONE, INJECTABLE	vial	8	222,720.00		RPS
89.	PROGESTERONE (BOVINE) ELISA KIT	kit	3	145,650.00		RPS
90.	PROSTAGLANDIN F2a ANALOGUE	dose	6,000	1,170,000.00		CALF
91.	PROSTAGLANDIN F2a ANALOGUE	dose	300	58,500.00		NDBH
92.	PROSTAGLANDIN F2a ANALOGUE	dose	30	5,850.00		RPS
93.	TRYPANOCIDAL DRUG (Anti-Surra Drug)	dose	2,350	1,762,500.00		BES
94.	TRYPANOCIDAL DRUG (Anti-Surra Drug)	dose	1,050	945,000.00		BES

TOTAL ABC: P 11,531,914.00

Section VII. Technical Specifications

Notes for Preparing the Technical Specifications

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

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

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

Sample Clause: Equivalency of Standards and Codes

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

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

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

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

In case of Renewal of Regular and Recurring Services, the Procuring Entity must indicate here the technical requirements for the service provider, which must include the set criteria in the conduct of its performance evaluation.

Technical Specifications

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

1.	**AI ALPHA SHEATH** - specific transparent dome head, dual-lateral delivery, unsplit, two lateral exit holes, crystalline plastic material - 50 pcs/pack	
2.	**AI GLOVES** - Shoulder Length at least 90cm (length), - Polyethylene gloves, - 25 microns (thickness), - 100 pcs/box	
3.	**AI GUN** - Universal - stainless steel	
4.	**AI STRAW SHEATHS** - 4.5mm outer diameter, 443mm in length, - made with medical grade non-toxic pvc material - 50 pcs/pack - Should fit Universal AI Gun (with split)	
5.	**ARTIFICIAL VAGINA FOR CATTLE** - Complete set Inclusions in set: - AV Casing with valve, pump and inner liner - 30cm -Robust tubular rubber casing with rounded openings. Valve assembly allows air to be added for adjustment of inner pressure.	
6.	**ARTIFICIAL VAGINA FOR GOAT** - Complete set Inclusions in set: - AV Casing with valve, pump and inner lner - 20cm -Robust tubular rubber casing with rounded openings. Valve assembly allows air to be added for adjustment of inner pressure.	
7.	**BULL HOLDER WITH CHAIN** - have pliers action and are used for short duration; - the chain keeps the grip closed; - assist handler to control animal with minimal risk of injury	
8.	**BULL RING APPLICATOR** - 3" diameter	

9.	**CIDAR APPLICATOR** - Must be compatible with CIDR EAZI-Breed	
10.	**CRYO GRIPPER FOR CRYOTANKS** - Size: Extends reach up to 32 inches with a strong leaf -style jaw - Suitable for Liquid Nitrogen - Use for retrieving Goblets and canes submerged in LN2	
11.	**DRENCHING GUN** - non-automatic - 60ml	
12.	**FORCEP** - at least 9-inch stainless steel	
13.	**GOBLET** - small (fit for Field tank) - Capacity: 100 x 0.5 ml or 210 x 0.25 ml straws - 25pcs/bag	
14.	**MILK DIPPER** - Nickel plated brass - with valve - sterilizable - Capacity: 40ml	
15.	**ORAL CANNULA / ORAL DRENCHER** - Fits any brand of Reusable Syringe via the Leur Lock System	
16.	**STAINLESS TONG** - 72 cm long	
17.	**STRAW CUTTER** - scissor type, - scissors intended for Artificial Insemination straw cutter scissor	
18.	**EAR TAG SUPPLIES** **-5000 pcs Animal Ear tag- Extra Large (XL)** Specification: with corresponding black male button-blank; Dimension: 3" W x 4.5/8" H; Material: High Grade Polyurethane; With infecta-Guard TM Coating/Treated Studs; Packaging: 25 pairs/bag; Color: Yellow; **- 2500 pcs Animal Ear tag- Large (L)**	

	Specifications: With corresponding black male button-blank; Dimension: 2 1/2" W x 2 3/4" H; Material: High Grade Polyurethane; With infecta-Guard TM Coating/Treated Studs; Packaging: 25 pairs/bag; Color: (Orange and Yellow) **-25 pcs Eartag Cutter** Material: Stainless Steel Product Dimensions: 7.01"L x 2.01"W **- 50 pcs Eartag Pen** Color Black Uneasily Remove or wash 5 years Guaranteed With Broad pump tip (until ink flows) Vapor Harmful, Flammable **- 50 pcs Eartagger replacement pin** Made of Metal; Must fit to the applicator destron fearing ProGripTM; Replaceable **- 30 pcs Eartagger** Tamper evident and electronic tags; Wide clip opening for insertion of tags in applicator; thicker "pro grip" handle coating for more comfort; same wide jaw for proper tag application; durable aluminum cast construction; strong spring return; tapered pin for easy release * Must have passed Trial Test **Note: Fits for ProGrip II Universal Tag Applicator** **- 1000 pcs Animal Rumen Bolus** Specifications:Dimension: 20.0mm +/-0.5 x 66.5mm +/-1.0 (ØxL); Weight: 72g; Material: Ceramic capsule Tecnology: HDX/FDX-B; Working frequency: 134.2 kHz; Certifications: ISO 11784/5 ICAR Approved; Reading Distance: Up to 28cm/25cm (HDX/FDX-B) with GES35 Reader, Up to 100 cm/85cm (HDX/FDX-B) with F310 Reader; * Must have passed Trial Test * Must be compatible to Datamars Bolus Applicator	
19.	**ALBENDAZOLE 15%** - 1 L/bot - Dosage: Maximum of 1ml/10 kg body wt	

	- Withdrawal period: - Meat: maximum of 14 days/ - Milk: maximum of 3 - 4 days - Minimum shelf life: 16 months from date of delivery - Must have passed trial test by end user	
20.	**ALBENDAZOLE 15%** - 1 L/bot - Dosage: Maximum of 1ml/10 kg body wt - Withdrawal period: - Meat: maximum of 14 days/ - Milk: maximum of 3 - 4 days - Minimum shelf life: 18 months from date of delivery - Must have passed trial test by end user	
21.	**ALBENDAZOLE 15%** - 1 L/bot - Dosage: Maximum of 1ml/10 kg body wt - Withdrawal period: - Meat: maximum of 14 days/ - Milk: maximum of 3 - 4 days - Minimum shelf life: 18 months from date of delivery - Must have passed trial test by end user	
22.	**AMINO ACID INJECTABLES / DEXTROLYTE** - 500 ml/bot - Minimum shelf life: 18 months from delivery date - Must have passed trial test by end user	
23.	**ATROPINE SULFATE** - 15mg 50ml/ bottle - Minimum shelf life: 24 months from date of delivery	
24.	**CALCIUM BOROGLUCONATE (CBG)** - 100ml - Minimum Shelf Life: 18 months from date of delivery - Must have passed trial test by end user	
25.	**CALIFORNIA MASTITIS TEST REAGENT** - CMT Liquid Concentrate (1 CMT concentrate = 1 gallon of working solution) - 1 gallon diluted reagent = approximately 350 tests - Minimum shelf life: 16 months from date of	

	delivery - Must have passed trial test by end user	
26.	**CAUSTIC SODA FLAKES** - 98-99% - 25kg/ pack - Minimum shelf life: 20 - 24 months from date of delivery	
27.	**CEFTIOFUR HCL/ SODIUM INJECTABLE** - 100 ml/bot - 50mg/ml - Zero withdrawal period - Minimum shelf life: 16 months from delivery date - Must have passed trial test by end user	
28.	**CEFTIOFUR HCL/ SODIUM INJECTABLE** - 100 ml/bot - 50mg/ml - Zero withdrawal period - Minimum shelf life: 18 months from delivery date - Must have passed trial test by end user	
29.	**CHLORTETRACYCLINE HYDROXIDE WOUND SPRAY** - 250 ml - active ingredient Chlortetracycline HCl 20mg per container - Cuteneous Suspension spray - Minimum shelf life: 16 months from date of delivery - Must have passed trial test by end user	
30.	**CHLORTETRACYCLINE HYDROXIDE WOUND SPRAY** - 250 ml - active ingredient Chlortetracycline HCl 20mg per container - Cuteneous Suspension spray - Minimum shelf life: 16 months from date of delivery - Must have passed trial test by end user	
31.	**DCM** - Dextrose 145 mg, calcium borogluonate180 mg, magnesium gluconate 62.1 mg - 500 ml - Minimum shelf life: 18 months from delivery date - Must have passed trial test by end user	

32.	**DCM** - Dextrose 145 mg, calcium borogluonate180 mg, magnesium gluconate 62.1 mg - 500 ml - Minimum shelf life: 18 months from delivery date - Must have passed trial test by end user	
33.	**DEXAMETHASONE SODIUM PHOSPHATE** - 2mg/ml - 100ml/bot - Minimum shelf life: at least 16 months from delivery date - Must have passed trial test by end user	
34.	**DEXAMETHASONE SODIUM PHOSPHATE** - 2mg/ml - 100ml/bot - Minimum shelf life: at least 18 months from delivery date - Must have passed trial test by end user	
35.	**DISINFECTANT** - 5 liters/container - Each liter must contain: Glutaraldehyde: 15%, Benzalkonium Chloride: 10%, Non-ionic surfactant: 1.5% - Active Ingredients: Glutaraldehyde and Quaternary Ammonium - Minimum shelf life: 18 months from date of delivery	
36.	**DORAMECTIN INJECTABLE** - 50ml/bot - Minimum shelf life: at least 16 months from delivery date - Must have passed trial test by end user	
37.	**DOXYCYCLINE** - 100 ml - Minimum shelf life: at least 16 months from delivery date - Must have passed trial test by end user	
38.	**EPINEPHRINE HYDROCHLORIDE** - 1ml/ampule - box of 50 - Minimum Shelf Life: 18 months from date of delivery	

39.	**FERRIC HYDROIDE + VITAMIN B 12 10%, INJECTABLE** - 100ml/bot - Must have passed trial test by end user - Minimum shelf life: 18 months from date of delivery	
40.	**HEMORRHAGIC SEPTECEMIA VACCINE** - 2ml/dose IM - Must have passed trial test by end user - Minimum Shelf Life: at least 18 months from date of delivery - Inclusive of: 20 pcs gel coolant	
41.	**HEPARIN SOLUTION, RX** - 10vial x 5ml - 5000 IU/ml - Solution for (IV/SC) injection - Anticoagulant	
42.	**INTRAMMAMMARY ANTIBIOTIC INFUSION TUBES FOR DRY ANIMALS** - 9g Composition (Each 9g syringe contains) :Cloxacillin benzathine: 1g - 24 syringes per carton - Minimum shelf life: At least 24 months from date of delivery - Must have passed trial test by the end user	
43.	**INTRAMMAMMARY ANTIBIOTIC INFUSION TUBES FOR LACTATING ANIMALS** - 10g Composition (each 10g syringe contains) : Kanamycin sulphate: 50mg, rocaine benzyl penicillin: 100,000 IU - 24 syringes per carton - Minimum shelf life: At least 24 months from date of delivery - Must have passed trial test by the end user	
44.	**IVERMECTIN INJECTABLE** - 50 ml/bot - content: Ivermectin 10 mg - Minimum shelf life: 16 months from delivery date - Must have passed trial test by end user	
45.	**IVERMECTIN INJECTABLE** - 50 ml/bot - content: Ivermectin 10 mg	

	- Minimum shelf life: 18 months from delivery date - Must have passed trial test by end user	
46.	**IVERMECTIN INJECTABLE** - 50 ml/bot - content: Ivermectin 10 mg - Minimum shelf life: 18 months from delivery date - Must have passed trial test by end user - Must attach BAI Clearance or FDA Registration Certificate	
47.	**KETOPROFEN** - 100mg/ml - Solution for injection (IM, IV) - Anti-inflammatory/ analgesic/ anti-pyretic) (NSAID) - Minimum shelf life: 18 months from date of delivery - Must have passed trial test by end user	
48.	**KETOPROFEN** - 100mg/ml - Solution for injection (IM, IV) - Anti-inflammatory/ analgesic/ anti-pyretic) (NSAID) - Minimum shelf life: 18 months from date of delivery - Must have passed trial test by end user	
49.	**L-CARNITINE USP,** - 200 mg - for cattle and buffaloes - Dosage: 5-6mg/kg body wt. for 3-5 days - for IM/IV use, - presentation: 30ml - 5 vials/box - Minimum shelf life: 12 months from delivery date - Must have passed trial test by end user	
50.	**LACTATED RINGER'S SOLUTION, RX** - 1000ml - Electrolytes in 1000ml: sodium 130 mmol, potassium 4 mmol, calcium 1.4 mmol, chloride 109 mmol, lactate 28 mmol - solution for IV infusion - Electrolyte replenisher - Sterile, non-pyrogenic - Single dose container	

51.	**LIDOCAINE HYDROCHORIDE 2%** - 20mg/ml - 50ml/bottle - Minimum shelf life: 24 months from date of delivery	
52.	**LIVER TONIC** - Choline chloride 2000mg, Protein Hydrolysate 50mg, Yeast Extract 40mg, Inositol 35mg, Niacin 24 mg, DL Panthenol 2.5 mg, Vitamin B12 3.3 mcg - 1 liter/ bot - Must have passed trial test by end user - Minimum shelf life: 18 months from date of delivery	
53.	**LIVER TONIC** - Choline chloride 2000mg, Protein Hydrolysate 50mg, Yeast Extract 40mg, Inositol 35mg, Niacin 24 mg, DL Panthenol 2.5 mg, Vitamin B12 3.3 mcg - 1 liter/ bot - Must have passed trial test by end user - Minimum shelf life: 18 months from date of delivery	
54.	**LIVER TONIC** - Choline chloride 2000mg, Protein Hydrolysate 50mg, Yeast Extract 40mg, Inositol 35mg, Niacin 24 mg, DL Panthenol 2.5 mg, Vitamin B12 3.3 mcg - 1 liter/ bot - Must have passed trial test by end user - Minimum shelf life: 18 months from date of delivery	
55.	**MINERAL BLOCK** - 5-7 kg. per block at - 4 blocks per box - Should contain the ff. composition: Sodium 38% Calcium 2.5% Magnesium 0.1% Manganese 1000 ppm min Copper 2400 ppm min Cobalt 50 ppm min Zinc 1000 ppm min Iron 3000 ppm min Iodine 100 ppm min Selenium 10 ppm min	
56.	**ORAL MULTIVITAMINS** - 1 liter	

	- Nutrient rich supplement - Minimum shelf life: 16 months from delivery date - Must have passed trial test by end user	
57.	**ORAL MULTIVITAMINS** - 1 liter - Nutrient rich supplement - Minimum shelf life: 16 months from delivery date - Must have passed trial test by end user	
58.	**OXYTETRACYCLINE LA 20%** - 100 ml/bot - Minimum shelf life: 16 months from delivery date - Must have passed trial test by end user	
59.	**OXYTETRACYCLINE LA 20%** - 100 ml/bot - Minimum shelf life: 16 months from delivery date - Must have passed trial test by end user	
60.	**PENTASULFATE SOLUTION** - 1L/bot - Minimum shelf life: 16 months from date of delivery - Must have passed trial test by end user	
61.	**PERMETHRIN SCREWWORM AEROSOL** - 10 ounce/can - Active ingredient: Permethrin: 0.5% - Topical wound spray - Minimum shelf life: 18 months from date of delivery - Must have passed trial test by end user	
62.	**POVIDONE IODINE (10% solution)** - 4l/gallon - Minimum shelf life: 24 months from delivery date - Must have passed trial test by end user	
63.	**SEALING POWDER** - orange - polyvinyl chloride powder - 750g	
64.	**TEAT SEALANT** - Composition: Each tube contains bismuth subnitrate (65% w/w) in an oil base, which is safe for use in non-lactating dairy cows	

	- For intramammary infusion - Packaging: 144 x 4g syringes	
65.	**TOLTAZURIL** - anti-coccidial - 250 ml/bot - Minimum shelf life: 24 months from delivery date - Must have passed trial test by end user	
66.	**VITAMIN ADE INJECTABLE** - Packaging: 100ml per bot. - Max of 4ml/ dose - Vitamin A 500,000 IU, D3 75,000 IU, Vit E 50 mg. - Must have passed trial test by end user - Minimum shelf life: 18 months from date of delivery	
67.	**VITAMIN ADE INJECTABLE** - Packaging: 100ml per bot. - Max of 4ml/ dose - Vitamin A 500,000 IU, D3 75,000 IU, Vit E 50 mg. - Must have passed trial test by end user - Minimum shelf life: 18 months from date of delivery	
68.	**VITAMIN ADE INJECTABLE** - Packaging: 100ml per bot. - Max of 4ml/ dose - Vitamin A 500,000 IU, D3 75,000 IU, Vit E 50 mg. - Must have passed trial test by end user - Minimum shelf life: 18 months from date of delivery	
69.	**VITAMIN ADE INJECTABLE** - Packaging: 100ml per bot. - Max of 4ml/ dose - Vitamin A 500,000 IU, D3 75,000 IU, Vit E 50 mg. - Must have passed trial test by end user - Minimum shelf life: 18 months from date of delivery	
70.	**VITAMIN B COMPLEX WITH LIVER EXTRACT** - 100 ml - Must have passed trial test by end user - Minimum shelf life: 18 months from date of delivery	
71.	**VITAMIN B12 BUTAPHOSPHAN , INJECTABLE** - 100ml/bot - Must have passed trial test by end user	

	- Minimum shelf life: 16 months from date of delivery	
72.	**VITAMIN B12 BUTAPHOSPHAN , INJECTABLE** - 100ml/bot - Must have passed trial test by end user - Minimum shelf life: 18 months from date of delivery	
73.	**AVIAN PPD TUBERCULIN ANTIGEN** - Each ml of tuberculin should contain: - Protein derivative of Mycobacterium avium subsp avium , strain D4 ER, 25,000 IU/ml - Dosage: 0.1 ml/dose - Packaging: 5ml/bot - Minimum Shelf Life: at least 12 months from date of delivery - Must have passed trial test by the end user	
74.	**BOVINE PPD TUBERCULIN ANTIGEN** - Each ml of tuberculin should contain: - Mycobacterium bovis, of at least 20,000 IU/ml - Dosage: 0.1 ml/dose - Packaging: 5ml/bot - Must have passed trial test by the end user Terms of delivery: 2-time progressive delivery **1st delivery:** 2000 doses * Minimum Shelf Life: at least 12 months from date of delivery **2nd delivery:** 1500 doses * Minimum Shelf Life: at least 12 months from date of delivery * Delivery period is 8 months after 1st delivery	
75.	**BOVINE PPD TUBERCULIN ANTIGEN** - Each ml of tuberculin should contain: - Mycobacterium bovis, of at least 20,000 IU/ml - Dosage: 0.1 ml/dose - Packaging: 5ml/bot - Must have passed trial test by the end user - Minimum Shelf Life: at least 12 months from date of delivery	
76.	**BRUCELLA TEST ANTIGEN** - 10mL bot equivalent to 330 reactions	

	- Rose Bengal stained acidified buffered antigen for rapid slide agglutination. - Composed of concentrated suspension of Brucella abortus (Weybridge strain 99) inactivated by heat and 0.5% phenol in an acid - Minimum shelf life: At least 16 months from date of delivery - Must have passed trail test by the end user - Inclusive of: 20 pcs gel coolant	
77.	**BRUCELLA TEST ANTIGEN** - 10mL bot equivalent to 330 reactions - Rose Bengal stained acidified buffered antigen for rapid slide agglutination. - Composed of concentrated suspension of Brucella abortus (Weybridge strain 99) inactivated by heat and 0.5% phenol in an acid - Minimum shelf life: At least 16 months from date of delivery - Must have passed trail test by the end user - Inclusive of: 20 pcs gel coolant	
78.	**CIDAR** - 1.38 grams progesterone implant - 10 pcs/pack - Minimum shelf life: 12 months from date of delivery must have passed trial test by the end user - Must have passed trial test by the end user - A certificate should be provided from the manufacturer that the supplier is authorized to sell the abovementioned biologics/hormone/drugs	
79.	**FASCIOLICIDAL DRUG** - 1 liter/bot - 100 mg/ml suspension - Active Ingredient : Triclabendazole - Dosage: Maximum of 1ml/10 kg. b.wt. - Withdrawal period: - Meat: Maximum of 42 days - Must have passed trial test by end user - Minimum shelf life: 24 monthS from date of delivery	
80.	**FASCIOLICIDAL DRUG** - 1 liter/bot - 100 mg/ml suspension - Active Ingredient : Triclabendazole - Dosage: Maximum of 1ml/10 kg. b.wt.	

	- Withdrawal period: - Meat: Maximum of 42 days - Must have passed trial test by end user - Minimum shelf life: 24 monthS from date of delivery	
81.	**GONADOTROPIN RELEASING HORMONE** - containing 43 mcg of Gonadorelin per ml - for 10 doses - 20 ml per bot - Minimum shelf life: 12 months from date of delivery - Must have passed trial test by the end user - A certificate should be provided from the manufacturer that the supplier is authorized to sell the abovementioned biologics/hormone/drugs	
82.	**GONADOTROPIN RELEASING HORMONE** - containing 43 mcg of Gonadorelin per ml - for 10 doses - 20 ml per bot - Minimum shelf life: 18 months from date of delivery - Must have passed trial test by the end user - A certificate should be provided from the manufacturer that the supplier is authorized to sell the abovementioned biologics/hormone/drugs	
83.	**HEMORRHAGIC SEPTECEMIA VACCINE** - 2ml/dose IM - Must have passed trial test by end user - Minimum Shelf Life: at least 16 months from date of delivery - Inclusive of: 20 pcs gel coolant Terms of delivery: 2-time progressive delivery **1st delivery:** 2000 doses **2nd delivery:** 2000 doses * Delivery period is 8 months after 1st delivery	
84.	**HORMONE, HUMAN CHORIONIC GONADOTROPIN (hCG)** - 1000 IU/vial - 5 vials/box	

	- 10,000 units hCG/single dose - 10 ml/vial - A certificate should be provided from the manufacturer that the supplier is authorized to sell the abovementioned biologics/hormone/drugs - Minimum shelf life: 12 months from date of delivery - Must have passed trial test by the end use	
85.	**HORMONE, HUMAN CHORIONIC GONADOTROPIN (hCG)** - 1000 IU/vial - 5 vials/box - 10,000 units hCG/single dose - 10 ml/vial - A certificate should be provided from the manufacturer that the supplier is authorized to sell the abovementioned biologics/hormone/drugs - Minimum shelf life: 12 months from date of delivery - Must have passed trial test by the end use	
86.	**PAG BLOOD KIT** - Bovine Pregnancy Test kit - 5x96 wells/box (for cattle, buffalo, goat or sheep) - type of samples: plasma or serum - Plate was coated with Pregnancy associated glycoproteins (PAG) - Minimum Shelf life: not less than 12 months from date of delivery to PCC - Must have passed trial test by end user	
87.	**PAG MILK KIT** - Bovine Pregnancy Test kit - 5x96 wells/box (for cattle, buffalo, goat or sheep) - type of sample: milk - Plate was coated with Pregnancy associated glycoproteins (PAG) - Minimum Shelf life: not less than 12 months from date of delivery to PCC - Must have passed trial test by end user	
88.	**PORCINE PITUITARY-DERIVED FOLLICLE STIMULATING HORMONE, INJECTABLE** - 20ml/vial - Minimum shelf life: 12 months from date of delivery - Must have passed trial test by the end user	

	- A certificate should be provided from the manufacturer that the supplier is authorized to sell the abovementioned biologics/hormone/drugs	
89.	**PROGESTERONE (BOVINE) ELISA KIT** - 96 tests - Quantitative determination of progesterone in Bovine milk or serum/plasma samples (Quantitative Protocol/Semiquantitative Protocol) - Competitive immunoassay - Minimum Shelf life: not less than 12 months from date of delivery	
90.	**PROSTAGLANDIN F2a ANALOGUE** - Dosage and route of administration: 2ml for intramuscular use in Carabao/Cattle - The drug has already been tested in local carabaos - Withdrawal period in milk: zero - FDA registration with revalidation certificate (if any) in case there are changes in Technical Information of the Product. The drug can still be used for more than fourteen (14) days after opening - Minimum Shelf life: 12 months from date of delivery - Must have passed trial test by the end-user - A certificate should be provided from the manufacturer that the supplier is authorized to sell the abovementioned biologics/hormone/drugs	
91.	**PROSTAGLANDIN F2a ANALOGUE** - Dosage and route of administration: 2ml for intramuscular use in Carabao/Cattle - The drug has already been tested in local carabaos - Withdrawal period in milk: zero - FDA registration with revalidation certificate (if any) in case there are changes in Technical Information of the Product. The drug can still be used for more than fourteen (14) days after opening - Minimum Shelf life: 12 months from date of delivery - Must have passed trial test by the end-user - A certificate should be provided from the manufacturer that the supplier is authorized to	

	sell the abovementioned biologics/hormone/drugs	
92.	**PROSTAGLANDIN F2a ANALOGUE** - Dosage and route of administration: 2ml for intramuscular use in Carabao/Cattle - The drug has already been tested in local carabaos - Withdrawal period in milk: zero - FDA registration with revalidation certificate (if any) in case there are changes in Technical Information of the Product. The drug can still be used for more than fourteen (14) days after opening - Minimum Shelf life: 12 months from date of delivery - Must have passed trial test by the end-user - A certificate should be provided from the manufacturer that the supplier is authorized to sell the abovementioned biologics/hormone/drugs	
93.	**TRYPANOCIDAL DRUG (Anti-Surra Drug)** - Isometamidium chloride hydrochloride,@400kg/dose - Route of administration: Intramuscular - Must provide diluent equivalent to the packaging of the drug - Withdrawal period: meat - 1 month; milk - none - Composition: 0.5 mg active ingredient / kg. body wt. of the drug - Minimum Concentration: 1 gram/50 ml. - Dosage Indication: Maximum of 1ml/20 kg. body wt. of the animal (e.g. maximum of 20ml/400kg buffalo per treatment) - Must have passed trial test by end user - Minimum shelf life: 18 months from date of delivery	
94.	**TRYPANOCIDAL DRUG (Anti-Surra Drug)** - Quinapyramine sulphate 1.5gm + Quinapyramine chloride 1.0g - Dosage indication: 0.025ml / kg. body wt. of the animal, inclusion of sterile distilled water. - Route of administration: Intramuscular or subcutaneous - Withdrawal period: meat - 21 days; milk - 4 days - Must have passed trial test by end user	

	- Minimum shelf life: 18 months from date of delivery	
	OTHER REQUIREMENTS: **a) TRIAL TEST** - Items to be offered must have been tested and passed trial test by the end user prior to the submission of bids (if required). - Certification duly signed by the end-user should be attached in the Schedule of Requirements/Technical Specifications. **b) MINIMUM SHELF LIFE** - Must be at least 12 months from date of delivery or acceptance made by PCC unless otherwise specified by the end-user	

CERTIFICATE OF TRIAL TEST BY END-USER
(attach this form in the Schedule of Requirements/Technical Specifications)

PROJECT:_____

SUPPLIER:_____

I/We hereby certify that the product/s offered has been used and considered acceptable/tested and passed trial test by the end-user.

ITEM NO	QTY	UNIT	ITEM AND DESCRIPTION	ENDUSER	OFFERED BRAND	NAME AND SIGNATURE OF EVALUATOR
18	1	lot	**EAR TAG SUPPLIES** **-5000 pcs Animal Eartag- Extra Large (XL)** Specification: with corresponding black male button-blank; Dimension: 3" W x 4.5/8" H; Material: High Grade Polyurethane; With infecta-Guard TM Coating/Treated Studs; Packaging: 25 pairs/bag; Color: Yellow; **- 2500 pcs Animal Eartag- Large (L)** Specifications: With corresponding black male button-blank; Dimension: 2 1/2" W x 2 3/4" H; Material: High Grade Polyurethane; With infecta-Guard TM Coating/Treated Studs; Packaging: 25 pairs/bag; Color: (Orange and Yellow) **-25 pcs Eartag Cutter** Material: Stainless Steel Product Dimensions: 7.01"L x 2.01"W **- 50 pcs Eartag Pen** Color Black Uneasily Remove or wash 5 years Guaranteed With Broad pump tip (until ink	ABGS		

			flows) Vapor Harmful, Flammable **- 50 pcs Eartagger replacement pin** Made of Metal; Must fit to the applicator destron fearing ProGripTM; Replaceable **- 30 pcs Eartagger** Tamper evident and electronic tags; Wide clip opening for insertion of tags in applicator; thicker "pro grip" handle coating for more comfort; same wide jaw for proper tag application; durable aluminum cast construction; strong spring return; tapered pin for easy release * Must have passed Trial Test **Note: Fits for ProGrip II Universal Tag Applicator** **- 1000 pcs Animal Rumen Bolus** Specifications:Dimension: 20.0mm +/-0.5 x 66.5mm +/-1.0 (ØxL); Weight: 72g; Material: Ceramic capsule Tecnology: HDX/FDX-B; Working frequency: 134.2 kHz; Certifications: ISO 11784/5 ICAR Approved; Reading Distance: Up to 28cm/25cm (HDX/FDX-B) with GES35 Reader, Up to 100 cm/85cm (HDX/FDX-B) with F310 Reader; * Must have passed Trial Test * Must be compatible to Datamars Bolus Applicator			
19	50	bot	**ALBENDAZOLE 15%** - 1 L/bot - Dosage: Maximum of 1ml/10 kg body wt - Withdrawal period: - Meat: maximum of 14 days/ - Milk: maximum of 3 - 4 days - Minimum shelf life: 16 months from date of delivery	BES		

			- Must have passed trial test by end user			
20	100	bot	**ALBENDAZOLE 15%** - 1 L/bot - Dosage: Maximum of 1ml/10 kg body wt - Withdrawal period: - Meat: maximum of 14 days/ - Milk: maximum of 3 - 4 days - Minimum shelf life: 18 months from date of delivery - Must have passed trial test by end user	CALF		
21	190	bot	**ALBENDAZOLE 15%** - 1 L/bot - Dosage: Maximum of 1ml/10 kg body wt - Withdrawal period: - Meat: maximum of 14 days/ - Milk: maximum of 3 - 4 days - Minimum shelf life: 18 months from date of delivery - Must have passed trial test by end user	NDBH		
22	20	bot	**AMINO ACID INJECTABLES / DEXTROLYTE** - 500 ml/bot - Minimum shelf life: 18 months from delivery date - Must have passed trial test by end user	LBRAF		
24	20	bot	**CALCIUM BOROGLUCONATE (CBG)** - 100ml - Minimum Shelf Life: 18 months from date of delivery - Must have passed trial test by end user	LBRAF		
25	20	bot	**CALIFORNIA MASTITIS TEST REAGENT** - CMT Liquid Concentrate (1 CMT concentrate = 1 gallon of working solution) - 1 gallon diluted reagent = approximately 350 tests - Minimum shelf life: 16 months from date of delivery	BES		

			- Must have passed trial test by end user			
27	15	bot	**CEFTIOFUR HCL/ SODIUM INJECTABLE** - 100 ml/bot - 50mg/ml - Zero withdrawal period - Minimum shelf life: 16 months from delivery date - Must have passed trial test by end user	BES		
28	40	bot	**CEFTIOFUR HCL/ SODIUM INJECTABLE** - 100 ml/bot - 50mg/ml - Zero withdrawal period - Minimum shelf life: 18 months from delivery date - Must have passed trial test by end user	NDBH		
29	10	can	**CHLORTETRACYCLINE HYDROXIDE WOUND SPRAY** - 250 ml - active ingradient Chlortetracycline HCl 20mg per container - Cuteneous Suspension spray - Minimum shelf life: 16 months from date of delivery - Must have passed trial test by end user	BES		
30	7	can	**CHLORTETRACYCLINE HYDROXIDE WOUND SPRAY** - 250 ml - active ingradient Chlortetracycline HCl 20mg per container - Cuteneous Suspension spray - Minimum shelf life: 16 months from date of delivery - Must have passed trial test by end user	LBRAF		
31	30	bot	**DCM** - Dextrose 145 mg, calcium borogluonate180 mg,magnesium gluconate 62.1 mg - 500 ml - Minimum shelf life: 18 months	LBRAF		

			from delivery date - Must have passed trial test by end user			
32	30	bot	**DCM** - Dextrose 145 mg, calcium borogluonate180 mg,magnesium gluconate 62.1 mg - 500 ml - Minimum shelf life: 18 months from delivery date - Must have passed trial test by end user	NDBH		
33	20	bot	**DEXAMETHASONE SODIUM PHOSPHATE** - 2mg/ml - 100ml/bot - Minimum shelf life: at least 16 months from delivery date - Must have passed trial test by end user	BES		
34	30	bot	**DEXAMETHASONE SODIUM PHOSPHATE** - 2mg/ml - 100ml/bot - Minimum shelf life: at least 18 months from delivery date - Must have passed trial test by end user	NDBH		
36	10	bot	**DORAMECTIN INJECTABLE** - 50ml/bot - Minimum shelf life: at least 16 months from delivery date - Must have passed trial test by end user	BES		
37	10	bot	**DOXYCYCLINE** - 100 ml - Minimum shelf life: at least 16 months from delivery date - Must have passed trial test by end user	BES		
39	20	bot	**FERRIC HYDROIDE + VITAMIN B 12 10%, INJECTABLE** - 100ml/bot - Must have passed trial test by end user - Minimum shelf life: 18 months from date of delivery	LBRAF		

40	650	dose	**HEMORRHAGIC SEPTECEMIA VACCINE** - 2ml/dose IM - Must have passed trial test by end user - Minimum Shelf Life: at least 18 months from date of delivery - Inclusive of: 20 pcs gel coolant	NDBH		
42	1	carton	**INTRAMMAMMARY ANTIBIOTIC INFUSION TUBES FOR DRY ANIMALS** - 9g Composition (Each 9g syringe contains) :Cloxacillin benzathine: 1g - 24 syringes per carton - Minimum shelf life: At least 24 months from date of delivery - Must have passed trial test by the end user	BES		
43	1	carton	**INTRAMMAMMARY ANTIBIOTIC INFUSION TUBES FOR LACTATING ANIMALS** - 10g Composition (each 10g syringe contains) : Kanamycin sulphate: 50mg, rocaine benzyl penicillin: 100,000 IU - 24 syringes per carton - Minimum shelf life: At least 24 months from date of delivery - Must have passed trial test by the end user	BES		
44	10	bot	**IVERMECTIN INJECTABLE** - 50 ml/bot - content: Ivermectin 10 mg - Minimum shelf life: 16 months from delivery date - Must have passed trial test by end user	BES		
45	7	bot	**IVERMECTIN INJECTABLE** - 50 ml/bot - content: Ivermectin 10 mg - Minimum shelf life: 18 months from delivery date - Must have passed trial test by end user	NDBH		

46	150	bot	**IVERMECTIN INJECTABLE** - 50 ml/bot - content: Ivermectin 10 mg - Minimum shelf life: 18 months from delivery date - Must have passed trial test by end user - Must attach BAI Clearance or FDA Registration Certificate	CALF		
47	15	bot	**KETOPROFEN** - 100mg/ml - Solution for injection (IM, IV) - Anti-inflammatory/ analgesic/ anti-pyretic) (NSAID) - Minimum shelf life: 18 months from date of delivery - Must have passed trial test by end user	LBRAF		
48	40	bot	**KETOPROFEN** - 100mg/ml - Solution for injection (IM, IV) - Anti-inflammatory/ analgesic/ anti-pyretic) (NSAID) - Minimum shelf life: 18 months from date of delivery - Must have passed trial test by end user	NDBH		
49	8	box	**L-CARNITINE USP,** - 200 mg - for cattle and buffaloes - Dosage: 5-6mg/kg body wt. for 3-5 days - for IM/IV use, - presentation: 30ml - 5 vials/box - Minimum shelf life: 12 months from delivery date - Must have passed trial test by end user	RPS		
52	10	bot	**LIVER TONIC** - Choline chloride 2000mg, Protein Hydrolysate 50mg, Yeast Extract 40mg, Inositol 35mg, Niacin 24 mg, DL Panthenol 2.5 mg, Vitamin B12 3.3 mcg - 1 liter/ bot - Must have passed trial test by end user	BES		

			- Minimum shelf life: 18 months from date of delivery			
53	10	bot	**LIVER TONIC** - Choline chloride 2000mg, Protein Hydrolysate 50mg, Yeast Extract 40mg, Inositol 35mg, Niacin 24 mg, DL Panthenol 2.5 mg, Vitamin B12 3.3 mcg - 1 liter/ bot - Must have passed trial test by end user - Minimum shelf life: 18 months from date of delivery	LBRAF		
54	30	bot	**LIVER TONIC** - Choline chloride 2000mg, Protein Hydrolysate 50mg, Yeast Extract 40mg, Inositol 35mg, Niacin 24 mg, DL Panthenol 2.5 mg, Vitamin B12 3.3 mcg - 1 liter/ bot - Must have passed trial test by end user - Minimum shelf life: 18 months from date of delivery	NDBH		
56	10	bot	**ORAL MULTIVITAMINS** - 1 liter - Nutrient rich supplement - Minimum shelf life: 16 months from delivery date - Must have passed trial test by end user	BES		
57	20	bot	**ORAL MULTIVITAMINS** - 1 liter - Nutrient rich supplement - Minimum shelf life: 16 months from delivery date - Must have passed trial test by end user	LBRAF		
58	10	bot	**OXYTETRACYCLINE LA 20%** - 100 ml/bot - Minimum shelf life: 16 months from delivery date - Must have passed trial test by end user	BES		
59	35	bot	**OXYTETRACYCLINE LA 20%** - 100 ml/bot - Minimum shelf life: 16 months from delivery date	NDBH		

			- Must have passed trial test by end user			
60	10	bot	**PENTASULFATE SOLUTION** - 1L/bot - Minimum shelf life: 16 months from date of delivery - Must have passed trial test by end user	BES		
61	12	can	**PERMETHRIN SCREWWORM AEROSOL** - 10 ounce/can - Active ingredient: Permethrin: 0.5% - Topical wound spray - Minimum shelf life: 18 months from date of delivery - Must have passed trial test by end user	NDBH		
62	5	gallon	**POVIDONE IODINE (10% solution)** - 4l/gallon - Minimum shelf life: 24 months from delivery date - Must have passed trial test by end user	NDBH		
65	10	bot	**TOLTAZURIL** - anti-coccidial - 250 ml/bot - Minimum shelf life: 24 months from delivery date - Must have passed trial test by end user	BES		
66	100	bot	**VITAMIN ADE INJECTABLE** - Packaging: 100ml per bot. - Max of 4ml/ dose - Vitamin A 500,000 IU, D3 75,000 IU, Vit E 50 mg. - Must have passed trial test by end user - Minimum shelf life: 18 months from date of delivery	BES		
67	135	bot	**VITAMIN ADE INJECTABLE** - Packaging: 100ml per bot. - Max of 4ml/ dose - Vitamin A 500,000 IU, D3 75,000 IU, Vit E 50 mg. - Must have passed trial test by end user	CALF		

			- Minimum shelf life: 18 months from date of delivery			
68	35	bot	**VITAMIN ADE INJECTABLE** - Packaging: 100ml per bot. - Max of 4ml/ dose - Vitamin A 500,000 IU, D3 75,000 IU, Vit E 50 mg. - Must have passed trial test by end user - Minimum shelf life: 18 months from date of delivery	LBRAF		
69	105	bot	**VITAMIN ADE INJECTABLE** - Packaging: 100ml per bot. - Max of 4ml/ dose - Vitamin A 500,000 IU, D3 75,000 IU, Vit E 50 mg. - Must have passed trial test by end user - Minimum shelf life: 18 months from date of delivery	NDBH		
70	60	bot	**VITAMIN B COMPLEX WITH LIVER EXTRACT** - 100 ml - Must have passed trial test by end user - Minimum shelf life: 18 months from date of delivery	NDBH		
71	60	bot	**VITAMIN B12 BUTAPHOSPHAN , INJECTABLE** - 100ml/bot - Must have passed trial test by end user - Minimum shelf life: 16 months from date of delivery	BES		
72	30	bot	**VITAMIN B12 BUTAPHOSPHAN , INJECTABLE** - 100ml/bot - Must have passed trial test by end user - Minimum shelf life: 18 months from date of delivery	LBRAF		
73	200	dose	**AVIAN PPD TUBERCULIN ANTIGEN** - Each ml of tuberculin should contain: - Protein derivative of Mycobacterium avium subsp avium , strain D4 ER, 25,000 IU/ml	BES		

			- Dosage: 0.1 ml/dose - Packaging: 5ml/bot - Minimum Shelf Life: at least 12 months from date of delivery - Must have passed trial test by the end user			
74	3,500	dose	**BOVINE PPD TUBERCULIN ANTIGEN** - Each ml of tuberculin should contain: - Mycobacterium bovis, of at least 20,000 IU/ml - Dosage: 0.1 ml/dose - Packaging: 5ml/bot - Must have passed trial test by the end user Terms of delivery: 2-time progressive delivery **<u>1st delivery:</u>** 2000 doses * Minimum Shelf Life: at least 12 months from date of delivery **<u>2nd delivery:</u>** 1500 doses * Minimum Shelf Life: at least 12 months from date of delivery * Delivery period is 8 months after 1st delivery	BES		
75	470	dose	**BOVINE PPD TUBERCULIN ANTIGEN** - Each ml of tuberculin should contain: - Mycobacterium bovis, of at least 20,000 IU/ml - Dosage: 0.1 ml/dose - Packaging: 5ml/bot - Must have passed trial test by the end user - Minimum Shelf Life: at least 12 months from date of delivery	NDBH		
76	10	bot	**BRUCELLA TEST ANTIGEN** - 10mL bot equivalent to 330 reactions - Rose Bengal stained acidified buffered antigen for rapid slide agglutination.	BES		

			- Composed of concentrated suspension of Brucella abortus (Weybridge strain 99) inactivated by heat and 0.5% phenol in an acid - Minimum shelf life: At least 16 months from date of delivery - Must have passed trial test by the end user - Inclusive of: 20 pcs gel coolant			
77	2	bot	**BRUCELLA TEST ANTIGEN** - 10mL bot equivalent to 330 reactions - Rose Bengal stained acidified buffered antigen for rapid slide agglutination. - Composed of concentrated suspension of Brucella abortus (Weybridge strain 99) inactivated by heat and 0.5% phenol in an acid - Minimum shelf life: At least 16 months from date of delivery - Must have passed trial test by the end user - Inclusive of: 20 pcs gel coolant	NDBH		
78	25	pack	**CIDAR** - 1.38 grams progesterone implant - 10 pcs/pack - Minimum shelf life: 12 months from date of delivery must have passed trial test by the end user - Must have passed trial test by the end user - A certificate should be provided from the manufacturer that the supplier is authorized to sell the abovementioned biologics/hormone/drugs	RPS		
79	25	bot	**FASCIOLICIDAL DRUG** - 1 liter/bot - 100 mg/ml suspension - Active Ingredient : Triclabendazole - Dosage: Maximum of 1ml/10 kg. b.wt. - Withdrawal period: - Meat: Maximum of 42 days - Must have passed trial test by end user	BES		

			- Minimum shelf life: 24 monthS from date of delivery			
80	20	bot	**FASCIOLICIDAL DRUG** - 1 liter/bot - 100 mg/ml suspension - Active Ingredient : Triclabendazole - Dosage: Maximum of 1ml/10 kg. b.wt. - Withdrawal period: - Meat: Maximum of 42 days - Must have passed trial test by end user - Minimum shelf life: 24 monthS from date of delivery	LBRAF		
81	22	vial	**GONADOTROPIN RELEASING HORMONE** - containing 43 mcg of Gonadorelin per ml - for 10 doses - 20 ml per bot - Minimum shelf life: 12 months from date of delivery - Must have passed trial test by the end user - A certificate should be provided from the manufacturer that the supplier is authorized to sell the abovementioned biologics/hormone/drugs	RPS		
82	50	vial	**GONADOTROPIN RELEASING HORMONE** - containing 43 mcg of Gonadorelin per ml - for 10 doses - 20 ml per bot - Minimum shelf life: 18 months from date of delivery - Must have passed trial test by the end user - A certificate should be provided from the manufacturer that the supplier is authorized to sell the abovementioned biologics/hormone/drugs	CALF		
83	5,000	dose	**HEMORRHAGIC SEPTECEMIA VACCINE** - 2ml/dose IM	BES		

			- Must have passed trial test by end user - Minimum Shelf Life: at least 16 months from date of delivery - Inclusive of: 20 pcs gel coolant Terms of delivery: 2-time progressive delivery **1st delivery:** 2000 doses **2nd delivery:** 2000 doses * Delivery period is 8 months after 1st delivery			
84	15	box	**HORMONE, HUMAN CHORIONIC GONADOTROPIN (hCG)** - 1000 IU/vial - 5 vials/box - 10,000 units hCG/single dose - 10 ml/vial - A certificate should be provided from the manufacturer that the supplier is authorized to sell the abovementioned biologics/hormone/drugs - Minimum shelf life: 12 months from date of delivery - Must have passed trial test by the end use	CALF		
85	12	box	**HORMONE, HUMAN CHORIONIC GONADOTROPIN (hCG)** - 1000 IU/vial - 5 vials/box - 10,000 units hCG/single dose - 10 ml/vial - A certificate should be provided from the manufacturer that the supplier is authorized to sell the abovementioned biologics/hormone/drugs - Minimum shelf life: 12 months from date of delivery - Must have passed trial test by the end use	RPS		
86	1	kit	**PAG BLOOD KIT** - Bovine Pregnancy Test kit - 5x96 wells/box (for cattle,	RPS		

			buffalo, goat or sheep) - type of samples: plasma or serum - Plate was coated with Pregnancy associated glycoproteins (PAG) - Minimum Shelf life: not less than 12 months from date of delivery to PCC - Must have passed trial test by end user			
87	1	kit	**PAG MILK KIT** - Bovine Pregnancy Test kit - 5x96 wells/box (for cattle, buffalo, goat or sheep) - type of sample: milk - Plate was coated with Pregnancy associated glycoproteins (PAG) - Minimum Shelf life: not less than 12 months from date of delivery to PCC - Must have passed trial test by end user	RPS		
88	8	vial	**PORCINE PITUITARY-DERIVED FOLLICLE STIMULATING HORMONE, INJECTABLE** - 20ml/vial - Minimum shelf life: 12 months from date of delivery - Must have passed trial test by the end user - A certificate should be provided from the manufacturer that the supplier is authorized to sell the abovementioned biologics/hormone/drugs	RPS		
90	6,000	dose	**PROSTAGLANDIN F2a ANALOGUE** - Dosage and route of administration: 2ml for intramuscular use in Carabao/Cattle - The drug has already been tested in local carabaos - Withdrawal period in milk: zero - FDA registration with revalidation certificate (if any) in case there are changes in Technical Information of the Product. The drug can still be used for more than fourteen (14) days after	CALF		

			opening - Minimum Shelf life: 12 months from date of delivery - Must have passed trial test by the end-user - A certificate should be provided from the manufacturer that the supplier is authorized to sell the abovementioned biologics/hormone/drugs			
91	300	dose	**PROSTAGLANDIN F2a ANALOGUE** - Dosage and route of administration: 2ml for intramuscular use in Carabao/Cattle - The drug has already been tested in local carabaos - Withdrawal period in milk: zero - FDA registration with revalidation certificate (if any) in case there are changes in Technical Information of the Product. The drug can still be used for more than fourteen (14) days after opening - Minimum Shelf life: 12 months from date of delivery - Must have passed trial test by the end-user - A certificate should be provided from the manufacturer that the supplier is authorized to sell the abovementioned biologics/hormone/drugs	NDBH		
92	30	dose	**PROSTAGLANDIN F2a ANALOGUE** - Dosage and route of administration: 2ml for intramuscular use in Carabao/Cattle - The drug has already been tested in local carabaos - Withdrawal period in milk: zero - FDA registration with revalidation certificate (if any) in case there are changes in Technical Information of the Product. The drug can still be used for more	RPS		

			than fourteen (14) days after opening - Minimum Shelf life: 12 months from date of delivery - Must have passed trial test by the end-user - A certificate should be provided from the manufacturer that the supplier is authorized to sell the abovementioned biologics/hormone/drugs			
93	2,350	dose	**TRYPANOCIDAL DRUG (Anti-Surra Drug)** - Isometamidium chloride hydrochloride,@400kg/dose - Route of administration: Intramuscular - Must provide diluent equivalent to the packaging of the drug - Withdrawal period: meat - 1 month; milk - none - Composition: 0.5 mg active ingredient / kg. body wt. of the drug - Minimum Concentration: 1 gram/50 ml. - Dosage Indication: Maximum of 1ml/20 kg. body wt. of the animal (e.g. maximum of 20ml/400kg buffalo per treatment) - Must have passed trial test by end user - Minimum shelf life: 18 months from date of delivery	BES		
94	1,050	dose	**TRYPANOCIDAL DRUG (Anti-Surra Drug)** - Quinapyramine sulphate 1.5gm + Quinapyramine chloride 1.0g - Dosage indication: 0.025ml / kg. body wt. of the animal, inclusion of sterile distilled water. - Route of administration: Intramuscular or subcutaneous - Withdrawal period: meat - 21 days; milk - 4 days - Must have passed trial test by end user - Minimum shelf life: 18 months from date of delivery	BES		

Note:
1. All signatories must be the unit head of unit/section/division concerned unless otherwise authorized by the unit head
2. For items with multiple end-users, any one of the end-user's signatures is accepted"

Ester B. Flores	ABGS	
Gabriel Alexis SP. Tubalinal	BES	
Edwin C. Atabay	CALF	
Lawrence P. Belotindos	LBRAF	
Cyril P. Balatzar	NDBH	
Excel Rio S. Maylem	RPS	

Our company hereby certify that the brand of products offered above are same products/goods with the same labelling and formulation.

Submitted by: _____

Name and signature of Supplier/Bidder: _____

Company Name: _____

Date: _____

Section VIII. Checklist of Technical and Financial Documents

Notes on the Checklist of Technical and Financial Documents

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

a. GPPB Resolution No. 09-2020 on the efficient procurement measures during a State of Calamity or other similar issuances that shall allow the use of alternate documents in lieu of the mandated requirements; or

b. Any subsequent GPPB issuances adjusting the documentary requirements after the effectivity of the adoption of the PBDs.

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

The Technical Component shall contain the following documents listed below. Kindly put tab markings/dog-ear codes on each of the requirements and arrange the documents in proper order.

Bidders are encouraged to submit the Technical and Financial Documents in three (3) copies for the following purposes:

1. **Original Copy** - reference of the BAC during the Opening of Bids/Evaluation and to be attached to the payment/voucher of the contractor/supplier
2. **Copy No. 1** - reference of the Technical Working Group for the Post-qualification
3. **Copy No. 2** - sealed copy to be provided to the Commission on Audit after the Opening of Bids

ANY discrepancy/insufficient or incomplete documents between the original and duplicate copies, the original shall prevail. Any document lacking in the bid envelope marked as ORIGINAL, the bid requirement will be marked as failed.

TECHNICAL COMPONENT (1st Envelope) please prepare in three (3) copies one original & two (2) duplicate copies (Copy No. 1 and Copy No. 2).

Refer to the illustration attached "Guide in proper sealing and marking of bids"

Checklist of Technical and Financial Documents

I. TECHNICAL COMPONENT ENVELOPE

Class "A" Documents

Legal Documents

☐ (a) Valid PhilGEPS Registration Certificate (Platinum Membership) (all pages) **in accordance with Section 8.5.2 of the IRR**;

Technical Documents

☐ (b) Statement of the prospective bidder of all its ongoing government and private contracts, including contracts awarded but not yet started, if any, whether similar or not similar in nature and complexity to the contract to be bid; **and**

☐ (c) Statement of the bidder's Single Largest Completed Contract (SLCC) similar to the contract to be bid, except under conditions provided for in Sections 23.4.1.3 and 23.4.2.4 of the 2016 revised IRR of RA No. 9184, within the relevant period as provided in the Bidding Documents; **and**

☐ (d) Original copy of Bid Security. If in the form of a Surety Bond, submit also a certification issued by the Insurance Commission **or** Original copy of Notarized Bid Securing Declaration; **and**

☐ (e) Conformity with the Technical Specifications, which may include production/delivery schedule, manpower requirements, and/or after-sales/parts, if applicable; **and**

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

Financial Documents

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

Class "B" Documents

☐ (h) If applicable, a duly signed joint venture agreement (JVA) in case the joint venture is already in existence **or** duly notarized statements from all the potential joint venture partners stating that they will enter into and abide by the provisions of the JVA in the instance that the bid is successful.

II. FINANCIAL COMPONENT ENVELOPE

☐ (i) Original of duly signed and accomplished Financial Bid Form; **and**

☐ (j) Original of duly signed and accomplished Price Schedule(s).

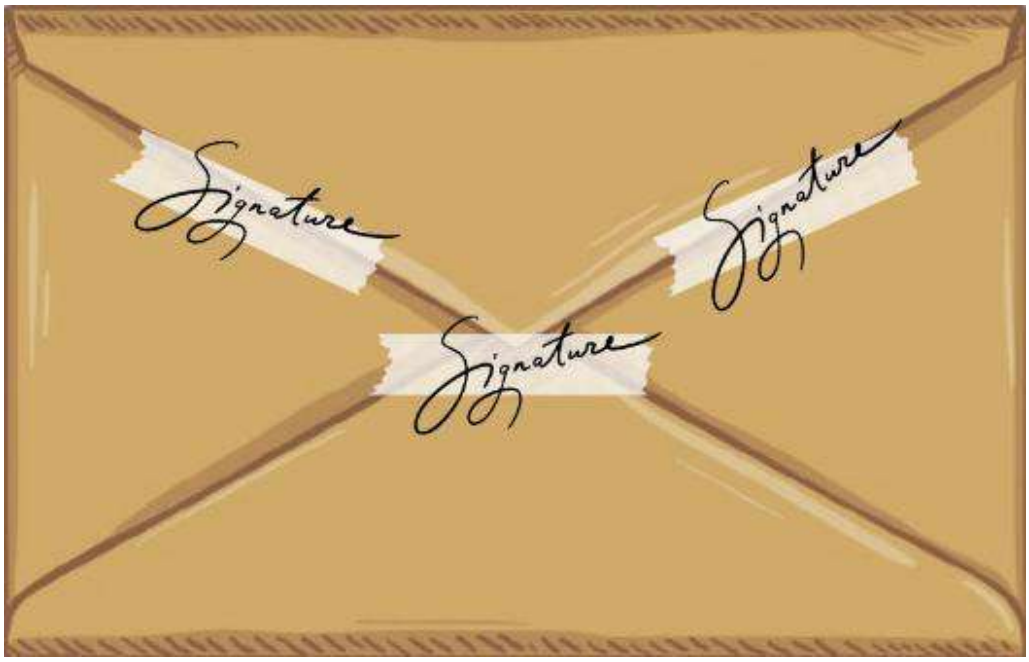
Other documentary requirements under RA No. 9184 (as applicable)

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

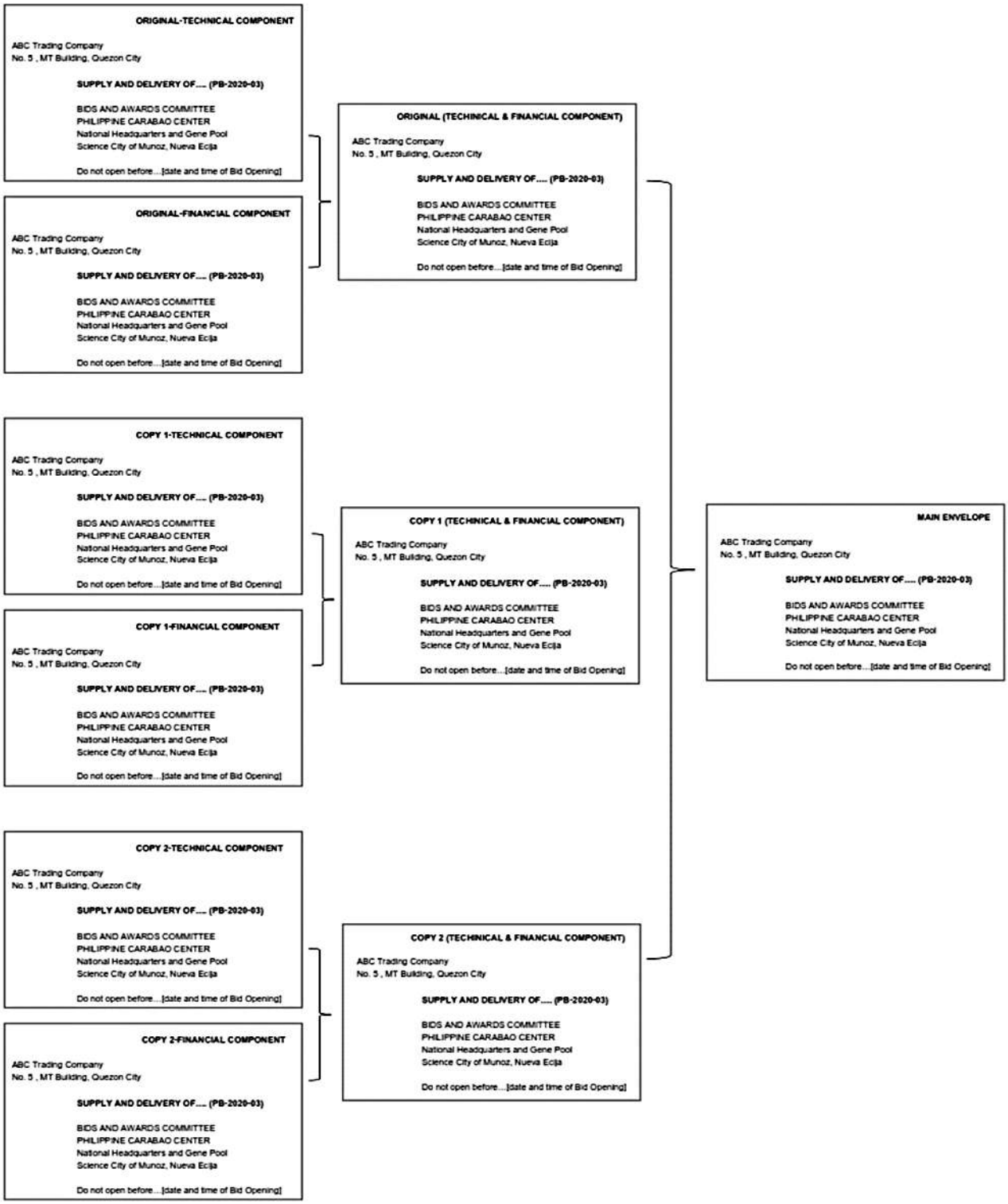
☐ (l) Certification from the DTI if the Bidder claims preference as a Domestic Bidder or Domestic Entity.

PROPER SEALING AND MARKING OF BIDS
(ALL 10 ENVELOPES)

- Signature should be at the **SEALED FLAP** of the envelope



PROPER SEALING AND MARKING OF BIDS
(ALL 10 ENVELOPES)



Other Instruction/s:

FOR GOODS AND INFRASTRUCTURE PROJECTS

1. All bidders shall upload and maintain in PhilGEPS a current and updated file of the following Class "A" eligibility documents under Sections 23.1(a) and 24.1(a) of RA 9184.

   Refer for PhilGEPS Advisory No. 2022-039 dated December 10, 2022

2. The notarized documents particularly the Bid Securing Declaration (BSD) and the Omnibus Sworn Statement (OSS) should have two different Document No. certified by the notary public, as these are two different or separate documents.

   All notarized documents should have a dry seal.

3. Declare all ongoing Government and Private Contracts (including contract awarded but not yet started, whether similar or not similar in nature.

FOR INFRASTRUCTURE PROJECTS

4. Do not add/delete items in the issued bid form/detailed estimates for infrastructure projects (modification of bid form).

5. Bidders are advised to distribute the discounts offered (if applicable) to the detailed estimates instead in the bid form.

6. The Certificate of Site Inspection should be signed by the PCC Center Director OR his/her authorized representative from the PCC. Prospective bidders may coordinate with the PCC regional centers being the end-user/project implementor. Electronic signature will be allowed for the said certificate.

Omnibus Sworn Statement (Revised)
[shall be submitted with the Bid]

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

AFFIDAVIT

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

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

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

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

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

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

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

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

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

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

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

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

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

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

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

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

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

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

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

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

Performance Securing Declaration (Revised)
[if used as an alternative performance security but it is not required to be submitted with the Bid, as it shall be submitted within ten (10) days after receiving the Notice of Award]

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

PERFORMANCE SECURING DECLARATION

Invitation to Bid: [Insert Reference Number indicated in the Bidding Documents]
To: [Insert name and address of the Procuring Entity]

I/We, the undersigned, declare that:

1. I/We understand that, according to your conditions, to guarantee the faithful performance by the supplier/distributor/manufacturer/contractor/consultant of its obligations under the Contract, I/we shall submit a Performance Securing Declaration within a maximum period of ten (10) calendar days from the receipt of the Notice of Award prior to the signing of the Contract.

2. I/We accept that: I/we will be automatically disqualified from bidding for any procurement contract with any procuring entity for a period of one (1) year for the first offense, or two (2) years **for the second offense**, upon receipt of your Blacklisting Order if I/We have violated my/our obligations under the Contract;

3. I/We understand that this Performance Securing Declaration shall cease to be valid upon:

    a. issuance by the Procuring Entity of the Certificate of Final Acceptance, subject to the following conditions:
        i. Procuring Entity has no claims filed against the contract awardee;
        ii. It has no claims for labor and materials filed against the contractor; and
        iii. Other terms of the contract; or

    b. replacement by the winning bidder of the submitted PSD with a performance security in any of the prescribed forms under Section 39.2 of the 2016 revised IRR of RA No. 9184 as required by the end-user.

IN WITNESS WHEREOF, I/We have hereunto set my/our hand/s this ____ day of [month] [year] at [place of execution].

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

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

Price Schedule for Goods Offered from Within the Philippines

[shall be submitted with the Bid if bidder is offering goods from within the Philippines]

For Goods Offered from Within the Philippines

Name of Bidder _____ Project ID No._____ Page ___of___

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

Name: _____

Legal Capacity: _____

Signature: _____

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

APPENDIX "1"

Bid Form for the Procurement of Goods

[shall be submitted with the Bid]

BID FORM

Date : _____

Project Identification No. : _____

To: [name and address of Procuring Entity]

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

If our Bid is accepted, we undertake:

a. to deliver the goods in accordance with the delivery schedule specified in the Schedule of Requirements of the Philippine Bidding Documents (PBDs);

b. to provide a performance security in the form, amounts, and within the times prescribed in the PBDs;

c. to abide by the Bid Validity Period specified in the PBDs and it shall remain binding upon us at any time before the expiration of that period.

[Insert this paragraph if Foreign-Assisted Project with the Development Partner:
Commissions or gratuities, if any, paid or to be paid by us to agents relating to this Bid, and to contract execution if we are awarded the contract, are listed below:

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

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

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

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

Name: _____

Legal capacity: _____

Signature: _____

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

Date: _____

Bid Securing Declaration Form

[shall be submitted with the Bid if bidder opts to provide this form of bid security]

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

BID SECURING DECLARATION
Project Identification No.: *[Insert number]*

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

I/We, 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 procurement 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 Declaration, within fifteen (15) days from receipt of the written demand by the procuring entity for the commission of acts resulting to the enforcement of the bid securing declaration under Sections 23.1(b), 34.2, 40.1 and 69.1, except 69.1(f),of the IRR of RA No. 9184; without prejudice to other legal action the government may undertake.

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; and
    c. I am/we are declared the bidder with the Lowest Calculated Responsive Bid, and I/we have furnished the performance security and signed the Contract.

IN WITNESS WHEREOF, I/We have hereunto set my/our hand/s this ____ day of *[month] [year]* at *[place of execution]*.

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

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