ANNEX F

Technical Specifications

Item	Specification	Statement of Compliance[1]
1.	**A. Type of Vaccine**: *Hepatitis B Vaccine* **Description**: *Recombinant Hepatitis B Vaccine 20mcg/ml Suspension for Injection (IM) -Single Dose* *Preservative Free* *1ml Vial* **Active Substances**: *Purified HBsAg 20mcg* **Dosage:** *1ml* (3 Doses) *1st Dose: at elected date* *2nd Dose: 1 month after the first dose* *3rd Dose: 6 months after the first dose* **Quantity:** *1,491 (497 pax x 3 doses)* **B. Description**: *HBsAg (Screen test for Hepatitis B prior the administration of vaccines)* **Quantity:** *497*	
2.	**Type of Vaccine:** *Tetanus Toxoid Vaccine* **Description:** *Tetanus Toxoid Adsorbed 40IU/0.5ml Suspension for IM Injection* **Active Substance:** *Tetanus Toxoid Adsorbed 40IU* ***Dosage:*** *0.5 ml.* **Quantity**: *489*	

[1] *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 Bidders 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 provisions of* ***ITB*** *Clause 3.1 (a)(ii) and/or GCC Clause 2.1 (a)(ii)..*

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.	Requirements: 1. Delivery of vaccines to DOT Head Office and Regional Offices. Supplier must prepare vaccines within 5 days upon receipt of Notice to Proceed document and must coordinate with the Head Office and Regional Offices regarding the HBsAg screen test, delivery and implementation schedule. 2. Cold Chain Storage during delivery must be maintained. 3. Vaccinators for the Head Office and Regional Offices will be provided by the supplier. They must be licensed nurses and should be under the supervision of a physician. For Regional Offices with a small number of employees, the physician will also be the vaccinator. Vaccinators and the physicians should follow the timeline of activities given below. 4. Supplier must submit the list of physician/s and nurse/s together with photocopies of their valid PRC licenses prior to the vaccination schedule. 5. Supplier must be FDA accredited. All vaccines should have Certificates of Product Registration from the FDA. 6. Expiration dates must not be less than 24 months from delivery date for Hepatitis B and Tetanus Toxoid vaccines. For vaccines which cannot meet this requirement, the supplier must submit a Certification Letter to the End User stating that these vaccines will automatically be replaced 4 months before the expiration dates. 7. Supplier must provide individual immunization record card for each employee. 8. All materials (refrigerator thermometer, etc.), medical supplies (alcohol, cotton balls, syringe, etc.), emergency medicines (Epinephrine, Diphenhydramine, etc.) and other	

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	necessary paraphernalia for the vaccination program shall be provided by the supplier. 9. Supplier is responsible for the disposal of all used materials and articles, especially the needles and syringes. These biological wastes should be disposed according to the procedures prescribed by the DOH Health Care Waste Management Manual. 10. Below is the recommended timeline of activities for the conduct of the Vaccination Program. The specific dates of immunization and screening test shall be agreed upon by the provider and the Head Office and Regional Offices.	
	Activity:	
	Screening Test (HBsAg) Date: within 5 days from receipt of Notice to Proceed	
	Vaccination Days Hepatitis Vaccination Day 1 (x 10 working days) – Hepatitis B Vaccine (1st Dose) Hepatitis Vaccination Day 2 (x10 working days – 1 month after 1st dose) – Hepatitis B Vaccine (2nd Dose) Anti-Tetanus Vaccination Day (x 7 working days – 1 month after the 2nd dose of Hepatitis B vaccine) – Tetanus Toxoid (One dose only) Hepatitis Vaccination Day 3 (x 10 working days – 6 months after the 1st dose) – Hepatitis B Vaccine (3rd Dose)	
	Submission of Vaccination Report Within 2 days after completion of vaccine administration	
	11. A series of meetings between the Medical Officer and the supplier must be done before, during and after the	

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	Vaccination Program, and midway in writing the Vaccination Report. 12. Vaccination Report should be submitted in hard and soft copy to the DOT Medical Clinic before the release of the Certificate of Completion. 13. Provider should submit a Vaccination Report based on the requirements of the DOT Medical Clinic. ***NOTE: Payment for the Hepatitis B vaccines will be based on the results of the screen test for Hepatitis B (HBsAg)***. ***ABC:*** ***Lot No. 1: PhP1,898,540.00 (Hepatitis B Vaccine)*** ***Lot No. 2: PhP100,245.00 (Tetanus Toxoid Vaccine)***	