

Section VII. Technical Specifications

Technical Specifications

Item	Specification	Statement of Compliance
		<i>[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.]</i>

ITEM NO.	ITEMS	SPECIFICATIONS	UNIT	STATEMENT OF COMPLIANCE
1	Face mask	<p>small size, good fit and good seal, N95 or its equivalent. With NIOSH marking and Testing Certificate Number marking on every mask or its equivalent. Must pass the standard test procedure. (Examples of acceptable disposable particulate respirators: Australia / New Zealand – P2, P3; China – II, I; European Union – FFP2, FFP3; Japan – 2nd class, 3rd class; Korea: 1st class, special; US – NIOSH certified N95, N99, N100).</p> <p>The brand being offered must have the following documents:</p>	piece	

		1. Food and Drug Administration (FDA) Certificate / Approval or Philippine Food and Drug Administration (FDA) Certificate of Medical Device Notification (CMDN) or Philippine Food and Drug Administration (FDA) Certificate of Exemption (COE) 2. Copy of Approval Letter / Accreditation Certificate or its equivalent 3. Copy of Test Report 4. Copy of Certificate of Authorized Distributorship		
2	Face mask	<p>regular size, good fit and good seal, N95 or its equivalent. With NIOSH marking and Testing Certificate Number marking on every mask or its equivalent. Must pass the standard test procedure. (Examples of acceptable disposable particulate respirators: Australia / New Zealand – P2, P3; China – II, I; European Union – FFP2, FFP3; Japan – 2nd class, 3rd class; Korea: 1st class, special; US – NIOSH certified N95, N99, N100).</p> <p>The brand being offered must have the following documents:</p> 1. Food and Drug Administration (FDA) Certificate / Approval or Philippine Food and Drug Administration (FDA) Certificate of Medical Device Notification (CMDN) or Philippine Food and Drug Administration (FDA) Certificate of Exemption (COE) 2. Copy of Approval Letter / Accreditation Certificate or its equivalent 3. Copy of Test Report 4. Copy of Certificate of Authorized Distributorship	piece	

TERMS & CONDITIONS:

1. The quantities specified are estimated quantities only and may be decreased based on the actual need of the hospital and availability of fund. It is understood that EAMC is not bound to purchase all of the quantities/items called for in the Invitation to Bid (IB).
2. Deliveries shall have at least eighteen (18) months expiration date/shelf-life. For items with shorter expiration date, the Supplier shall submit an assurance letter prior to deliver for approval of the Head of the Procuring Entity (HOPE) or his authorized representative.

<i>Conforme:</i>
<i>Company/Bidder's Name:</i>
<i>Name and Signature of Authorized Representative:</i>
<i>Date:</i>