

PRICE INFORMATION SCHEDULES**SCHEDULE 1: PRICE SUMMARY AND TERMS OF PAYMENT**

Tenderers shall set out in this Schedule:

1. **Price Summary** including the breakdown of the overall Tender Price stated in the Tender Form and any other terms of payment. The breakdown shall be detailed in the following format according to the summary of items and quantities required in the Tender. The price shall include any labour charges, custom tax and with holding tax.
2. Payment will be made locally and in Brunei Dollars. Payment shall be made in accordance with the provisions in the Contract.

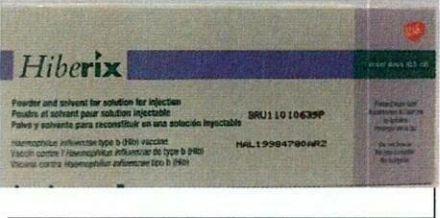
ITEM	DESCRIPTION	QTY	UNIT PRICE B\$	TOTAL PRICE B\$
1	PBL3055 HAEMOPHILUS INFLUENZAE TYPE B (HIB) VACCINE	150 UNITS		
2	PBL1557 INFANRIX IPV (4 IN 1) VACCINE	180 UNITS		
3	PBL1454 INFANRIX HEXA (6 IN 1) VACCINE	320 UNITS		
4	PBL1519 MEASLES, MUMPS, RUBELLA VACCINE (MMR)	230 UNITS		
5	PBL1200 PNEUMOCOCCAL POLYSACCHARIDE CONJUGATE (13-VALENT VACCINE)	210 UNITS		

NOTE:

- The Government reserves the right to give PARTIAL AWARD.
- The Government may accept the whole or any item(s) of tender.
- The Price quotes need to be maintained in items of partial award.
- **SPECIFICATIONS AND CATALOGUE MUST BE SUBMITTED AND CLEARLY LABELED.**



COMPLIANCE AGREEMENT TABLE FOR SUPPLIER

Serial	Item Name	HEAMOPHILUS INFLUENZAE TYPE B (HIB) VACCINE		
	Catalogue / Specification			
(a)	(b)	(c)	(d)	(e)
DESCRIPTION & USER REQUIREMENTS		Comply (√)	Not Comply (√)	Remark
1	Validity of offer price shall be at least 6 months from the closing date of submission of quote			
2	Sample of the actual product being offered in untampered original pack including package insert.(for controlled drugs and psychotropic drugs see item 3)			
3	<p>A CLEAR QUALITY PICTURE of the primary and secondary packaging of the product being offered: showing name/brand if item, strength and form/preparation, from all sides/angles. Each picture is to be printed in colour, and this document stamped with suppliers' tenderers' official stamp.</p> <p>Additionally, pictures of the following: For tablets/capsules a. Appearance of individual tablets/capsules b. If the item is in strip pack, the back and front of the strip .For injections: a. Appearance of individual vial/ampoule/syringe</p>			
4	Certificate of Analysis			
5	A copy of any of the following: - Product licence certificate - Log of submission for registration of the product			
6	Registration status in any of the accepted "benchmark country"			
7	Letter of authorization from the product Licence holder, where applicable			



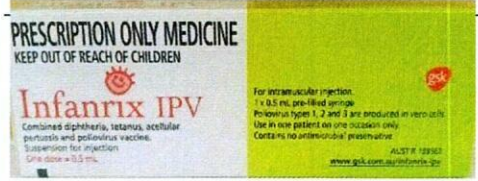
8	Justification on price increase if the same product has been previously supply to Ministry of defence from the same supplier/distributor			
9	Latest local content			
10	Product shelf-life information / Description - Shelf life period of time from the date of manufacture			
11	On delivery, goods should have a minimum expiry date of 24 months unless item has short expiry (eg: Vaccines) or agreed to be accepted by MRS. Any period less than 24 months shall provide Letter of Undertaking			
12	The storage labelling should be in accordance with ASEAN stability guideline and should be based on stability evaluation of the drug product. Terms such as "ambient condition", "room temperature" or "does not require any special storage conditions" should be avoid			
13	Description & Specification:			
	Haemophilus b conjugate vaccine (tetanus toxoid conjugate) is an active immunizing agent that is used to prevent infection caused by the Haemophilus influenza type b (Hib) bacteria. The vaccine works by causing your body to produce its own protection (antibodies) against the disease.			
14	Manufacture by Glaxo Smith Kline Pharmaceutical Sdn.Bhd.			
15	Type: Injection			
16	Packing size: 1 vial with lyophilised vaccine + 1 pre-filled with solvent + 2 needles			
17	Dose: 1 dose/0.5ml			
18	Country Origin :			
19	Quality Assurance:			
20	Delivery Period/Time :			

Note :

- a. Will cancel the product, if the delivery product is not same as specification given by Medical Supply Centre Bolkliah.
- b. Expired date must be written in the Delivery Order / Invoice.
- c. Any Brand should be okay, as long it fullfill as our specification requirements and Supplier MUST STATED the brand name of the product.



COMPLIANCE AGREEMENT TABLE FOR SUPPLIER

Serial	Item Name	INFANRIX IPV (4 in 1) VACCINE		
	Catalogue / Specification			
(a)	(b)	(c)	(d)	(e)
DESCRIPTION & USER REQUIREMENTS		Comply (√)	Not Comply (√)	Remark
1	Validity of offer price shall be at least 6 months from the closing date of submission of quote			
2	Sample of the actual product being offered in untampered original pack including package insert.(for controlled drugs and psychotropic drugs see item 3)			
3	<p>A CLEAR QUALITY PICTURE of the primary and secondary packaging of the product being offered: showing name/brand if item, strength and form/preparation, from all sides/angles. Each picture is to be printed in colour, and this document stamped with suppliers' tenderers official stamp.</p> <p>Additionally,pictures of the following:</p> <ul style="list-style-type: none"> - For tablets/capsules <ul style="list-style-type: none"> a.Appearance of individual tablets/capsules b.If the item is in strip pack,the back and front of the strip - For injections: <ul style="list-style-type: none"> a. Apperance of individual vial/ampoule/syringe 			
4	Certificate of Anaiysis			
5	<p>A copy of any of the following</p> <ul style="list-style-type: none"> - Product licence certificate - Log of submission for registration of the product 			
6	Registration status in any of the accepted "benchmark country"			
7	Letter of authorization from the product Licence holder, where applicable			



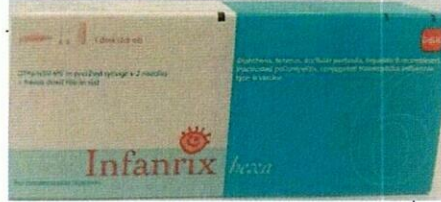
8	Justification on price increase if the same product has been previously supply to Ministry of defence from the same supplier/distributor			
9	Latest local content			
10	Product shelf-life information / Description - Shelf life period of time from the date of manufacture			
11	On delivery, goods should have a minimum expiry date of 24 months unless item has short expiry (eg: Vaccines) or agreed to be accepted by MRS. Any period less than 24 months shall provide Letter of Undertaking			
12	The storage labelling should be in accordance with ASEAN stability guideline and should be based on stability evaluation of the drug product. Terms such as "ambient condition", "room temperature" or "does not require any special storage conditions" should be avoid			
13	<u>Description & Specification:</u>			
	Infanrix IPV is a quadri-valent vaccine. It protects children from diphtheria, pertussis, tetanus, polio. It contains DTAP and inactivated polio vaccine.			
14	Manufacture by Glaxo Smith Kline Pharmaceutical Sdn.Bhd.			
15	Type: Injection			
16	Packing size: 1 x 0.5ml pre-filled injection			
17	Dose: 1 dose/0.5ml			
18	Country Origin :			
19	Quality Assurance:			
20	Delivery Period/Time :			

Note :

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- b. Expired date must be written in the Delivery Order / Invoice.
- c. Any Brand should be okay, as long it fullfill as our specification requirements and Supplier MUST STATED the brand name of the product.



COMPLIANCE AGREEMENT TABLE FOR SUPPLIER

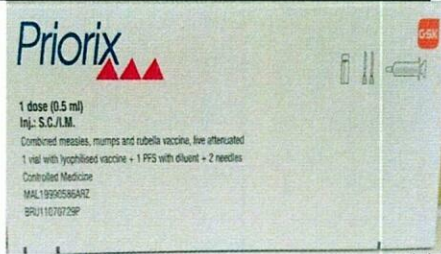
Serial	Item Name	INFANRIX HEXA (6 in 1) VACCINE		
	Catalogue / Specification			
(a)	(b)	(c)	(d)	(e)
DESCRIPTION & USER REQUIREMENTS		Comply (√)	Not Comply (√)	Remark
1	Validity of offer price shall be at least 6 months from the closing date of submission of quot			
2	Sample of the actual product being offered in untampered original pack including package insert.(for controlled drugs and psychotropic drugs see item 3)			
3	A CLEAR QUALITY PICTURE of the primary and secondary packaging of the product being offered:showing name/brand if item,strength and form/preparation, from all sides/angles. Each picture is to be printed in colour,and this document stamped with suppliers tenderers official stamp. Additionally,pictures of the following: -For tablets/capsules a.Appearance of individual tablets/capsules b.If the item is in strip pack,the back and front of the strip -For injections: a.Apperance of individual vial/ampoule/syringe			
4	Certificate of Anaiysis			
5	A copy of any of the following - Product licence certificate - Log of submission for registration of the product			
6	Registration status in any of the accepted "benchmark country"			



7	Letter of authorization from the product Licence holder, where applicable			
8	Justification on price increase if the same product has been previously supply to Ministry of defence from the same supplier/distributor			
9	Latest local content			
10	Product shelf-life information / Description - Shelf life period of time from the date of manufacture			
11	On delivery, goods should have a minimum expiry date of 24 months unless item has short expiry (eg: Vaccines) or agreed to be accepted by MRS. Any period less than 24 months shall provide Letter of Undertaking			
12	The storage labelling should be in accordance with ASEAN stability guideline and should be based on stability evaluation of the drug product. Terms such as "ambient condition", "room temperature" or "does not require any special storage conditions" should be avoid			
13	Description & Specification:			
	Infanrix hexa is a vaccine used to against diphtheria, tetanus, pertussis (whooping cough), hepatitis B, poliomyelitis (polio) and diseases such as bacterial meningitis caused by the bacterium Haemophilus influenzae type b (Hib).			
14	Manufacture by Glaxo Smith Kline Pharmaceutical Sdn.Bhd.			
15	Type: Injection			
16	Packing size: 1 vial and 1 pre-filled syringe + 2 needles . 1 vial - powder and pre filled syringe - suspension			
17	Dose: 1 dose/0.5ml			
18	Country Origin :			
19	Quality Assurance:			
20	Delivery Period/Time :			
<p>Note :</p> <p>a. Will cancel the product , if the delivery product is not same as specification given by Medical Supply Centre Bolkiah .</p> <p>b. Expired date must be write in the Delivery Order / Invoice.</p> <p>c. Any Brand should be okay , as long it fullfill as our specification requirements and Supplier MUST STATED the brand name of the product.</p>				



COMPLIANCE AGREEMENT TABLE FOR SUPPLIER

Serial	Item Name	MEASLES, MUMPS, RUBELLA VACCINE		
	Catalogue / Specification			
(a)	(b)	(c)	(d)	(e)
DESCRIPTION & USER REQUIREMENTS		Comply (✓)	Not Comply (✓)	Remark
1	Validity of offer price shall be at least 6 months from the closing date of submission of quot			
2	Sample of the actual product being offered in untampered original pack including package insert.(for controlled drugs and psychotropic drugs see item 3)			
3	<p>A CLEAR QUALITY PICTURE of the primary and secondary packaging of the product being offered:showing name/brand if item,strength and form/preparation, from all sides/angles. Each picture is to be printed in colour,and this document stamped with suppliers tenderers official stamp.</p> <p>Additionally,pictures of the following:</p> <p>-For tablets/capsules</p> <p>a.Appearance of individual tablets/capsules</p> <p>b.If the item is in strip pack,the back and front of the strip</p> <p>-For injections:</p> <p>a.Apperance of individual vial/ampoule/syringe</p>			
4	Certificate of Anaiysis			
5	<p>A copy of any of the following</p> <p>-Product licence certificate</p> <p>-Log of submission for registration of the product</p>			



6	Registration status in any of the accepted "benchmark country"			
7	Letter of authorization from the product Licence holder, where applicable			
8	Justification on price increase if the same product has been previously supply to Ministry of defence from the same supplier/distributor			
9	Latest local content			
10	Product shelf-life information / Description - Shelf life period of time from the date of manufacture			
11	On delivery, goods should have a minimum expiry date of 24 months unless item has short expiry (eg: Vaccines) or agreed to be accepted by MRS. Any period less than 24 months shall provide Letter of Undertaking			
12	The storage labelling should be in accordance with ASEAN stability guideline and should be based on stability evaluation of the drug product. Terms such as "ambient condition", "room temperature" or "does not require any special storage conditions" should be avoid			
13	Description & Specification:			
	MMR's vaccine is a vaccine indicated for active immunization for the prevention of measles, mumps, and rubella in individuals.			
14	Manufacture by Glaxo Smith Kline Pharmaceutical Sdn.Bhd.			
15	Type: Injection			
16	Packing size: 1 vial and 1 pre-filled syringe + 2 needles . 1 vial - powder and pre filled syringe - suspension			
17	Dose: 1 dose/0.5ml			
18	Country Origin :			
19	Quality Assurance:			
20	Delivery Period/Time :			

Note :

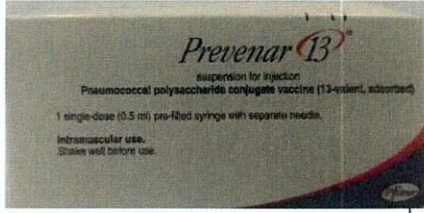
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COMPLIANCE AGREEMENT TABLE FOR SUPPLIER

Serial	Item Name	PNEUMOCOCCAL POLYSACCHARIDE CONJUGATE (13-VALENT VACCINE)		
	Catalogue / Specification			
(a)	(b)	(c)	(d)	(e)
DESCRIPTION & USER REQUIREMENTS		Comply (✓)	Not Comply (✓)	Remark
1	Validity of offer price shall be at least 6 months from the closing date of submission of quot			
2	Sample of the actual product being offered in untampered original pack including package insert. (for controlled drugs and psychotropic drugs see item 3)			
3	<p>A CLEAR QUALITY PICTURE of the primary and secondary packaging of the product being offered: showing name/brand if item, strength and form/preparation, from all sides/angles. Each picture is to be printed in colour, and this document stamped with suppliers' tenderers official stamp.</p> <p>Additionally, pictures of the following:</p> <ul style="list-style-type: none"> - For tablets/capsules <ol style="list-style-type: none"> a. Appearance of individual tablets/capsules b. If the item is in strip pack, the back and front of the strip - For injections: <ol style="list-style-type: none"> a. Appearance of individual vial/ampoule/syringe 			
4	Certificate of Analysis			
5	<p>A copy of any of the following</p> <ul style="list-style-type: none"> Product licence certificate Log of submission for registration of the product 			
6	Registration status in any of the accepted "benchmark country"			



7	Letter of authorization from the product Licence holder, where applicable			
8	Justification on price increase if the same product has been previously supply to Ministry of defence from the same supplier/distributor			
9	Latest local content			
10	Product shelf-life information / Description - Shelf life period of time from the date of manufacture			
11	On delivery, goods should have a minimum expiry date of 24 months unless item has short expiry (eg: Vaccines) or agreed to be accepted by MRS. Any period less than 24 months shall provide Letter of Undertaking			
12	The storage labelling should be in accordance with ASEAN stability guideline and should be based on stability evaluation of the drug product. Terms such as "ambient condition", "room temperature" or "does not require any special storage conditions" should be avoid			
13	Description & Specification:			
	PNEUMOCOCCAL POLYSACCHARIDE CONJUGATE (13-VALENT VACCINE) is used to prevent pneumonia, meningitis, and blood and ear infections.			
14	Type: Injection			
15	Paking size: 1 single dose (0.5ml) pre-filled syringe with separate needle			
16	Country Origin :			
17	Quality Assurance:			
18	Delivery Period/Time :			

Note :

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