

ANNEX A

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	PSY4891.L DIPHENYDRAMINE 14MG. AMMONIUM CI 135MG, CITRATE 57MG PER 5ML (100ML) & (3.8L)	400 GALLONS/2000 BOTTLES		
2	PSY0500.L TRIPROLIDINE HCI 1.25 MG + PSEUDOPHEDRINE 30MG PER 5ML (60ML) & (3.8L)	400 GALLONS/2000 BOTTLES		
3	PSY1224.L PARACETEMOL SUSPENSION 250MG/5ML (3.8L) & (100 ML)	400 GALLONS/2000 BOTTLES		
4	PSY0809.L CARBOCISTEINE SYRUP (SVMC) 250MG/5ML (100ML)	1000 BOTTLES		
5	PSY0712.L TRIPROLIDINE HCI 1.25MG, PSEUDOEPHEDRINE HCI 30MG, DEXTROMETHORPHAN HBRI 15MG PER 5ML SYRUP	1000 BOTTLES		
GRAND TOTAL B\$				

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.**



ANNEX B

COMPLIANCE AGREEMENT TABLE FOR SUPPLIER

Serial	Item Name	DIPHENYDRAMIDE 14mg, AMMONIUM Cl 135mg, CITRATE 57mg PER 5ml		
	Catalogue / Specification	As Attached		
(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 <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	Description & Specification: DIPHENYDRAMIDE 14mg, AMMONIUM CI 135mg, CITRATE 57mg PER 5ml is indicated as an expectorant for control of cough due to cold or allergy.			
14	Type: Syrup			
15	Paking size: 1 gallon equal to 3.8 litre			
16	Manufactured: Sunward Pharmaceutical Sdn.Bhd			
17	Country Origin :			
18	Quality Assurance:			
19	Delivery Period/Time :			
Note : <ol style="list-style-type: none"> Will cancel the product , if the delivery product is not same as specification given by Medical Supply Centre Bolkiah . Expired date must be write in the Delivery Order / Invoice. 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	TRIPROLIDINE HCl 1.25mg + PSEUDOPHEDRINE 30mg PER 5ml		
	Catalogue / Specification	As Attached		
(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: TRIPROLIDINE HCl 1.25mg + PSEUDOPHEDRINE 30mg per 5ml is indicated in nasal and respiratory congestion, common cold, acute sinusitis, and allergic rhinitis			
14	Type: Syrup			
15	Packing size: 1 gallon equal to 3.8 litre			
16	Manufactured: Sunward Pharmaceutical Sdn.Bhd			
17	Country Origin :			
18	Quality Assurance:			
19	Delivery Period/Time :			
Note : a. Will cancel the product , if the delivery product is not same as specification given by Medical Supply Centre Bolkiah . b. Expired date must be write 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	PARACETAMOL 250mg/5ml SUSPENSION		
	Catalogue / Specification	As Attached		
(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 <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	Description & Specification: PARACETAMOL 250mg/5ml SUSPENSION is for relief of pain and fever, influenza, rheumatic and joint pain, cramps and other muscular pains.			
14	Type: Suspension			
15	Paking size: 1 gallon equal to 3.8 litre			
16	Manufactured: Beacon Pharmaceuticals PTE.LTD			
17	Country Origin :			
18	Quality Assurance:			
19	Delivery Period/Time :			
Note : a. Will cancel the product , if the delivery product is not same as specification given by Medical Supply Centre Bolkiah . b. Expired date must be write 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	CARBOCISTEINE SYRUP (SCMC)		
	Catalogue / Specification	As Attached		
(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 <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	Description & Specification: CARBOCISTEINE SYRUP is for used in the treatment of problems in the airways when there is too much mucus or when the mucus is too sticky, including chronic obstructive airways disease (COPD).			
14	Type: Syrup			
15	Paking size: 1 bottle equal to 100ml			
16	Manufactured: Duopharma Manufacturing (Bangi) Sdn.Bhd			
17	Country Origin :			
18	Quality Assurance:			
19	Delivery Period/Time :			
Note : a. Will cancel the product , if the delivery product is not same as specification given by Medical Supply Centre Bolkiah . b. Expired date must be write 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	TRIPROLIDINE HCl 1.25mg, PSUEDOEPHEDRINE HCl 30mg, DEXTROMETHORPHAN HBr 15mg PER 5ml		
	Catalogue / Specification	As Attached		
(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: TRIPROLIDINE HCl 1.25mg, PSEUDOEPHEDRINE HCl 30mg, DEXTROMETHORPHAN HBr 15mg is indicated for the relief of dry irritating coughs such as those associated with common cold, upper respiratory tract infections, allergic (seasonal) rhinitis, vasomotor (perennial) rhinitis and hay fever.			
14	Type: Syrup			
15	Packing size: 1 bottle equal to 120ml			
16	Manufactured: Xepa-Soul Pattinson (Malaysia) Sdn Bhd			
17	Country Origin :			
18	Quality Assurance:			
19	Delivery Period/Time :			
Note : <ol style="list-style-type: none"> Will cancel the product , if the delivery product is not same as specification given by Medical Supply Centre Bolkiah . Expired date must be write in the Delivery Order / Invoice. Any Brand should be okay , as long it fullfill as our specification requirements and Supplier MUST STATED the brand name of the product. 				

