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 (3.8L)	300 GALLONS		
2	PSY0500.L TRIPROLIDINE HCI 1.25 MG + PSEUDOPHEDRINE 30MG PER 5ML (3.8L)	300 GALLONS		
3	PSY1224.L PARACETEMOL SUSPENSION 250MG/5ML (3.8L OR 4 LITRE)	300 GALLONS		
4	PSY0809.L CARBOCISTEINE SYRUP (SCMC)	600 BOTTLES		
5	PSY0712.L TRIPROLIDINE HCI 1.25MG, PSEUDOEPHEDRINE HCI 30MG, DEXTROMETHORPHAN HBRI 15MG PER 5ML	600 BOTTLES		
6	PSY1913.L CO-AMOXICLAV SUSPENSION 228ML/5ML	300 BOTTLES		
7	PSY7721.L AMOXYCILLIN SYRUP 125MG/5ML	400 BOTTLES		
8	PCT2391 ORAL REHYDRATION SALT 50 SACHETS PER BOX	1000 BOXES		

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**		
	Catalouge / 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 psychotripic 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 duideline 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 Cl 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 :

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 + PSEUDOPHEDRINE 30mg PER 5ml**		
	Catalouge / 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 psychotripic 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 duideline 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	**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 :

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**		
	Catalouge / 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 psychotripic 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 duideline 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)**		
	Catalouge / 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 psychotripic 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 duideline 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**		
	Catalouge / 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 psychotripic 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 duideline 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	**Paking 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 :

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	**CO-AMOXICLAV (AUGMENTIN) 228mg/5ml SUSPENSION**		
	Catalouge / 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 psychotripic 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 duideline 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:** **CO-AMOXICLAV (AUGMENTIN) 228mg/5ml SUSPENSION** is indicated for short-term treatment of bacterial infections such as upper respiratory tract infections, lower respiratory tract infections, skin and soft tissue infections, bone and joint infections, dental infections and other infections (septic abortion, puerperal sepsis and intra-abdominal sepsis).			
14	**Type:** Suspension			
15	**Paking size:** 1 bottle equal to 70ml			
16	**Manufactured:** Healal Pharmaceuticals 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	**AMOXYCILLIN 125mg/5ml SYRUP**		
	Catalouge / 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 psychotripic 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 duideline 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:** **AMOXYCILLIN 125mg/5ml SYRUP** is indicated for susceptible infections including ear,nose,throat (ENT), genitourinary tract, skin and skin structures, lower respiratory, acute uncomplicated gonorrhea.			
14	**Type:** Suspension			
15	**Paking size:** 1 bottle equal to 60ml			
16	**Manufactured:** Duopharma (M) 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	**ORAL REHYDRATION SALTS**		
	Catalouge / 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 psychotripic 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 duideline 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:** **ORAL REHYDRATION SALTS** is for replacement of water and electrolytes lost during moderate to severe diarrhoea.			
14	**Type:** Powder			
15	**Paking size:** 1 box : 50 sachets			
16	**Manufactured:** Pharmaniaga Manufacturing Berhad			
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.