The Health Sciences Center

# BIDS & AWARDS COMMITTEE IV (BAC IV)

Proj. Ref. No. BAC1-2022-08-0051-B

End-User Property and Supply Division Opening of Bids: 16 September 2022

Project Supply and Delivery of Drugs and Medicines - Tablet/Oral Solids

ITEM	QTY	UNIT	ITEM DESCRIPTION	ABC PER UNIT	QUOTATIONS (all taxes included)	
NO.	QII	CIVII	TIEM DESCRIPTION	(PHP)	in figures	in words
			TABLETS/ORAL SOLID			
1	9,040	piece	Aciclovir 400mg tablet	30.00		
2	1,200		Aripiprazole 10mg tablet	62.45		
3	45,541	piece	Aspirin 80mg entric coated tablet	2.10		
4	17,514		Atorvastatin calcium 10mg tablet	7.88		
5	23,939		Atorvastatin calcium 20mg tablet	4.20		
6	47,730		Atorvastatin calcium 40mg tablet	9.45		
7	22,610		Atorvastatin calcium 80mg tablet	8.93		
8	540		Betahistine (as hydrochloride/dihydrochloride) 8mg tablet	12.60		
9	8,060		Carbamazepine 200mg tablet	4.26		
10	32,661		Carvedilol 25mg tablet	5.25		
11	1,856		Cefixime 200mg capsule	8.40		
12	19,114		Cefuroxime axetil 500mg tablet	9.98		
13	2,600		Celecoxib 100mg capsule	2.10		
14	8,773	piece	Cetirizine di-Hydrochloride 10mg tablet	3.59		
15	2,440	piece	Clindamycin Hydrochloride 150mg capsule	6.80		
16	9,427	piece	Clonidine Hydrochloride 75mcg tablet	9.56		
17	36,769	piece	Clopidogrel 75mg tablet	6.04		
18	28,142	piece	Clozapine 100mg tablet	30.45		
19	7,110	piece	Co-Amoxiclav (amoxicillin + potassium clavulanate) 875 mg amoxicillin (as trihydrate) + 125 mg potassium clavulanate per tablet	7.19		
20	5,160	piece	Eperisone Hydrochloride 50mg tablet	29.00		
	,		Ferrous/folic acid 60 mg elemental iron/400 mcg per tablet/capsule/film			
21	30,340	piece	coated tablet	1.88		
22	38,550		Folic acid 5mg tablet/capsule	4.20	<b> </b>	
23	16,720		Isoniazid + Rifampicin 75 mg + 150 mg tablet	9.45		
24	10,165		Isosorbide-5-mononitrate 30mg MR tablet/capsule	6.83		
25	5,590		Isosorbide-5-mononitrate 60mg MR tablet/capsule	12.00	$\vdash$	
26	2,165		Lamotrigine 50mg tablet	5.64	$\vdash$	
27	582		Lamotrigine 100mg tablet	15.75	$\vdash$	
28	7,240		Levofloxacin 500mg tablet	7.35	$\vdash$	
29	7,937		Levofloxacin 750mg tablet	12.60		
30	3,040		Levothyroxine (as sodium/anhydrous sodium) 150mcg tablet	10.89		
31			Loratadine 10mg tablet/film-coated tablet	1.41	$\vdash$	
32		_	Mefenamic acid 500mg tablet/capsule	11.55		
33	60,002		Mercaptopurine 50mg tablet	25.00		
34	90,258		Metformin Hydrochloride 500mg tablet/film-coated tablet	0.51		
35	31,080		Methotrexate (as base or sodium salt) 2.5mg tablet	4.72		
36	8,619	_	Metoclopramide Hydrochloride 10mg tablet	6.30	<del>                                     </del>	
37	5,635		Metoprolol tartrate 100mg tablet	2.10	-	
38	18,905		Metoprolol tartrate 50mg tablet	1.05	-	
39	20,337		Omeprazole 20mg capsule	2.24	-	
40	26,520		Potassium Chloride 600 mg tablet	18.00		
41	27,049	_	Potassium citrate 10mEq tablet	7.35		
42	28,053		Prednisone 10mg tablet	4.40	-	
43	5,983		Quetiapine (as fumarate) 100mg tablet	88.88		
44	9,226		Quetiapine (as fumarate) 25mg tablet	12.60		
45	5,047	_	Quetiapine (as fumarate) 200mg tablet	70.00		
46	5,700		Ranitidine Hydrochloride 300mg tablet	3.80		
47	6,592		Rifaximin 200mg tablet	66.36		
48	8,010	piece	Risperidone 1mg tablet	10.89	<u> </u>	

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			·	-	
49	19,091	piece	Risperidone 2mg tablet	40.95	
50	10,745		Rosuvastatin calcium 10mg tablet	12.81	
51	14,910		Rosuvastatin calcium 20mg tablet	22.05	
52	14,423	101000	Sodium Valproate + Valproic Acid 500 mg (333 mg sodium valproate + 145	11.95	
32			mg valproic acid) controlled release tablet		
53	48,102	piece	Tamoxifen citrate 20mg tablet	2.15	
54	16,695	piece	Tamsulosin 400mcg prolonged release film-coated tablet	98.87	
55	2,960	piece	Tamsulosin Hydrochloride 200mcg capsule	28.00	
56	23,316	piece	Telmisartan 40mg tablet	16.87	
57	28,087	piece	Telmisartan 80mg tablet	28.13	
58	2,170		Tenofovir disoproxil fumarate 300 mg tablet	41.22	
59	1,810	piece	Tenofovir alafenamide fumarate 25 mg tablet	70.00	
60	2,312	piece	Topiramate 25mg tablet	26.00	
61	24,360	piece	Tramadol Hydrochloride 50mg capsule/dispersible tablet	7.35	
62	26,792	piece	Trimetazidine Hydrochloride 35mg tablet	11.55	
63	1,620		Ursodeoxycholic acid 250mg capsule	28.35	
64	700	piece	Valaciclovir (as hydrochloride) 500mg tablet	252.00	
65	1,050		Valsartan 80mg tablet/film-coated tablet	11.23	
66	73,879	piece	Vitamin B1 100 mg + B6 5 mg + B12 50 microgram per tablet/capsule	4.20	
	TOTAL APPROVED BUDGET FOR THE CONTRACT: PhP13,941,239.76				

Approved	by:
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Dean CHARLOTTE M. CHIONG, MD, PhD	,
Chairperson	

(Signature over Printed Name of Presid	dent/Gen. Manager)
(Name & Address of Company)	<u> </u>

## **TERMS AND CONDITIONS**

- 1. Indicate the brand and packing of the item/s offered.
- 2. The brand offered must be at least five (5) years commercially available in the market from date of opening of bids. Proof of this shall be the initial Certificate of Product Registration (CPR) issued by the Food and Drug Administration (FDA).
- 3. Submit the following documents, submission should be per product, with tab and per item number. Two (2) copies for the Valid Certificate of Product Registration and Certificate of Analysis (COA).
- 3.1. Memorandum of Agreement (MOA) and Certificate of exclusive/authorized distributorship between the manufacturer and distributors/suppliers must have certification from their principals that they are the exclusive distributor of the drug products authorized t tender for the product on behalf of the principal and that all commitments made by them shall be honored by the principal in ca termination of distributorship agreement.
- **3.2. Valid Certificate of Product Registration (CPR)** issued by the Food and Drug Administration (FDA).
- The name of the respective distributor should appear on the submitted CPR of the drug.

Note: CPRs that will expire within three (3) months from the date of opening of bids should present the Official Receipt of renewal of application v Document Tracking log for the CPR from the FDA.

**3.3. Certificate of Analysis (COA)** for the products offered (batch to be delivered if awarded) duly issued by an FDA-accredited laboratory (low and should contain information indicated in monograph of the drug. Sample analyzed must not be expired during the time of bith The result of assay submitted must be in the specific brand and should be in the exact dosage formulation of the drug being bidd

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In cases where local laboratories are unavailable to perform drug assays, assays done abroad is accepted. The local COA is prefer and given more weight in the evaluation and awarding process.

- **3.4.** A notarized certificate that it is the innovator drug (if applicable).
- 3.5. Certificate of Good Manufacturing Process (CGMP).
- 3.6. Valid License to Operate (LTO).
- 3.7. A notarized certificate that the offered brand has not been subject to product complaint/product recall for the past three (3) years.
- **3.8. Certificate of Acceptance** from at least three (3) major hospitals issued within the year and should be supported with Sales Invoice (for new item/brand offered only).
- 3.9. A notarized certificate that there are sufficient stocks for the offered item/s for one (1) year.
- 4. The offered drug must conform to the latest Philippine Food and Drug Administration (FDA) Approved by:
- 4.1. For all tablets and capsules
- 4.1.1. All tablets/capsules should be in foil or blister pack. A picture of the blister pack (front and back) should be submitted.
- 4.1.2. Each individual flap in the tablet or capsule blister pack should be labeled with the generic r Dean VICENTE Y. BELIZARIO, JR., M.D.
- 4.1.3. Dosage form and strength of the Active Pharmaceutical Ingredients (API) should appear on Chairperson
- **4.1.4.** Name of drug, lot or batch number and expiry date must appear on every standard blister pack/foil strip and on the container or inner packi However, if the product is not restricted for dispensing in quantities less than the standard blister pack or foil strip, the batch or lot number and expir should appear on each unit.
- **4.1.5.** Inner label must be the same as the outer label.
- **4.1.6.** A complete drug literature/product insert must accompany the product.
- 5. The following must be complied with specific for cytotoxic injectable drugs.
- **5.1. For cytotoxic injectable drugs, winning bidders are required to** *provide Material Safety Data Sheet (MSDS) and to submit Drug Profile* to the Pharmacy Department per company under the first Purchase Order.
- 5.2. Winning bidders for cytotoxic injectable drugs are required to provide at least three (3) spill kits per company under the first Purchase Ord
- 6. New brands offered shall be subject to further evaluation and shall require the following:
- **6.1.** Validation of the submitted Certificate of Acceptance from at least three (3) major hospitals;
- **6.2.** Justification from end-user/s to validate the acceptance of the good/s offered (to be facilitated by PGH-PSD).
- 7. For the supply and delivery of awarded drugs and medicines.
- **7.1.** Delivery of the goods is required as stated in the request of the end-user, commencing on the 3rd working day of notification through confirmal fax/email that the approved Call-off/ Notice to Supplier (NTS) is already available for pick up.
- **7.2.** Delivery schedule (whichever is applicable):
- **7.2.1.** within seven (7) calendar days;
- 7.2.2. as may be called for;
- 7.2.3. staggered delivery within three (3) months
- \* 50% of the total quantity within seven (7) calendar days and 25% each for the succeeding months

Note: The end-user has the right to adjust the quantity to be delivered depending on the actual need of the hospital

- 7.3. Deliveries shall have at least two (2) years expiration date.
- **7.4.** Delivery of goods **with product complaint shall be put on hold** until receipt of the final decision of the PGH management whether to product with the acceptance or to cancel/return the items.
- **7.5.** Delivered **items found to be non-formulary at any given time shall be returned** to the company and a credit memo shall be issued.
- 7.6. Stocks delivered are subject to random sampling for testing as to quality and conformity to label. Testing fee at supplier's expense.
- **7.7.** Stocks with lot #/batch different from the submitted Certificate of Analysis (COA) will be subjected to testing as to quality and conformity t Testing fee at supplier's expense.

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- **7.8.** All items that had been pulled out for various reasons, a credit memo shall be issued by the Contractor within one (1) month, otherwise, a de shall be processed by UP Manila PGH and the amount will be deducted from any amount due to Supplier.
- 7.9. It is understood that the Supplier is legally responsible to deliver all issued CALL-OFF/s (Purchase Order) and failure to deliver the first as scheduled shall mean automatic cancellation of the Call-Off and Notice to Execute Framework Agreement (NEFA). Purchase from other source for shall be effected immediately to provide the requirements of the hospital. Penalty to the defaulting contractor shall be charged accordingly.
- 8. Failure to comply with the submission of the required documents shall be ground for post-disqualification in accordance with RA 91
- 9. Compliance with RA 9184 and other applicable laws.

	Approved by:	
	Dean CHARLOTTE M. CHIONG, MD, PhD Chairperson	
(Signature over Printed Name of President/Gen. Manager)		
(Name & Address of Company)		