Republic of the Philippines
DEPARTMENT OF HEALTH
OFFICE OF THE SECRETARY

PURCHASE ORDER

Procurement of Gliclazide 30mg MR Tablet

IB No. 2018-075

Supplier:	**EuroCare Pharma Inc.**	PO No. :	**GOP-2018-04-032**
Address:	Km. 21 East Service Road, Cupang Muntinlupa City	Date:	MAY 31 2018
Telephone No.	Tel no: (02) 623-6721 Fax No. (02) 807-1057	Mode of Procurement:	Competitive Bidding
TIN	006-360-258-000		

Sir/Madam:

Please furnish this office of the following articles subject to the terms and conditions contained herein:

Place of Delivery:	DOH Warehouse(s) or Service Provider(s) in Metro Manila	Delivery Term:	
Date of Delivery:	Within Sixty (60) calendar days upon receipt of approved Notice to Proceed (NTP)	Mode of Payment:	**Upon Delivery and Acceptance**

Item No.	Unit	Description	Quantity	Unit Cost	Amount
1	Treatment Pack	**Gliclazide 30mg MR tablet** **Name of Manufacturer: IPCA Laboratories Ltd.** **Country of Origin:** India **Brand:** Glycinorm-MR 30 **Technical Specification:** • Oral • 30mg Modified Release (MR) tablet • Must be commercially available in the Philippine market for the past two (2) years **Upon delivery the following shall be complied with:** • **Shelf Life:** Shelf life of at least *twenty four (24) months* from the date of manufacture but not less than *eighteen (18) months* from the date of delivery. • **Packaging Instructions:** • Primary Packaging: Thirty (30) tablets per DOH Complete Treatment Pack • Secondary Packaging: Standard packaging of the manufacturer including product insert or encryption/imprint inside the box as approved by PFDA. • **Labeling Instructions:** Standard labeling instruction as approved by Philippine Food and Drug Administration pursuant to Administrative Order No. 2016-0008. **In addition to the labelling requirement of FDA:** • On each **blister pack/foil strip and treatment pack,** the following should be legibly imprinted or stickered with non-removable or permanent sticker/label that is binding and with residue and tearing, if removed: **Philippine Government Property-DOH** **Not for Sale** • On each **corrugated carton**, the following should be legibly imprinted or stickered with non-removable or permanent sticker/label that is binding and with residue and tearing, if removed: **Philippine Government Property-DOH** **Not for Sale** Manufacturing Date: _____ Expiration Date: _____ Batch/Lot No.: _____ • **Product Recall/Replacement** The supplier must ensure the quality of drugs and if there will be problems in the quality, the supplier will recall and replace the drugs distributed in the regions and health centers based on the Guidelines on Product Recall, FDA Circular No. 2016-012. **Other Requirements for Payment *(for Drugs Medicines):*** **1.** Within a period of thirty (30) days from receipt of delivery, a one hundred percent (100%) payment arrangement shall be made to the company upon receipt of satisfactory results of analysis from PFDA. **2.** If, however, by thirty (30) days from receipt of delivery, PFDA cannot realease the result of laboratory analysis, the company shall be paid fifty (50%) of the due amount. The remaining fifty percent (50%) balance is to be paid after obtaining a satisfactory PFDA report of analysis. **3.** The supplier shall be charged of the laboratory testing fee for the test analysis of the delivered drugs and medicines. The payment shall be made directly by the supplier to the PFDA Cashier's Office. Otherwise, the Laboratory test result will not be released to the Logistic Management Division (LMD) of the Department of Health. **The inspection and tests that will be conducted are:** 1. Upon delivery, the Goods shall undergo preliminary physical Inspection by the Inspection Team of the PROCURING ENTITY to ascertain the physical condition and acceptability of the Goods. 2. The supplier shall promptly replace the equivalent quantity of Goods taken as samples without cost to the PROCURING ENTITY. 3. Pending PFDA analysis, said products should not be distributed to end-users nor shall it be used until such time it is cleared by PFDA laboratory testing 4. If PFDA inspection or results of laboratory analysis show major violations, the entire product line of the drug is temporarily suspended from accreditation, regardless of the batch or lot in question. 5. All health commodities **with failed FDA test analysis shall immediately be pulled out by the suppliers from the DOH warehouse or from the warehouse of the DOH service provider within five (5) working days** upon receipt of FDA test result from the Logistic Management Division - Administrative Service (LMD-AS). Failure to comply within the prescribed time shall compel this Office to have the subject commodities pulled out by the third party logistics service provider of the DOH with **the hauling and freight fees chargeable against the concerned supplies.** In the observance of the above-mentioned timeline, please coordinate with the LMD-AS of the DOH. **Purpose**: For 2018 procurement for the Diabetes Mellitus Medicine Access Program Note: Subject to the conditions stated in the Bidding Documents	168,000	28.00	4,704,000.00
Four Million Seven Hundred Four Thousand Philippine Pesos					4,704,000.00

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Liquidated Damages (Section 68 of the Revised IRR of RA9184): All contracts executed in accordance with the Act and this IRR shall contain a provision on liquidated damages which shall be payable by the contractor in case of breach thereof. For the procurement of Goods, Infrastructure Projects and Consulting Services, the amount of the liquidated damages shall be at least equal to one-tenth of one percent (0.001) of the cost of the unperformed portion for every day of delay. Once the cumulative amount of liquidated damages reaches ten percent (10%) of the amount of the contract, the Procuring Entity may rescind or terminate the contract, without prejudice to other courses of action and remedies available under the circumstances.

Warranty (Section 62 of the Revised IRR of RA 9184): In order to assure that manufacturing defects shall be corrected by the supplier, a warranty security shall be required from the contract awardee for a minimum period of three (3) months, in the case of Expendable Supplies,or a minimum period of one (1) year, in the case of Non-expendable Supplies, after acceptance by the Procuring Entity of the delivered supplies.The obligation for the warranty shall be covered by either retention money in an amount equivalent to at least one percent (1%) of every progress payment, or a special bank guarantee equivalent to at least one percent (1%) of the total contract price. The said amounts shall only be released after the lapse of the warranty period or, in the case of Expendable Supplies, after consumption thereof: Provided, however,That the supplies delivered are free from patent and latent defects and all the conditions imposed under the contract have been fully met.

Conforme:

KHRISTINE S. CORTEZ
Signature Over Printed Name of Supplier

8/13/18
Date

By Authority of the Secretary of Health:

MYRNA C. CABOTAJE, MD, MPH, CESO III
Director IV
Disease Prevention and Control Bureau

Fund Cluster: _____

Funds Available: _____

LORICA C. RABAGO
OIC ACCOUNTING DIVISION

ORS/BURS No.: ~~03101101-2018-06-03971~~

Date of the ORS/BURS: 6-1-18

Amount: ₱ 4,704,000.00

Building 1, San Lazaro Compound, Rizal Avenue, Sta. Cruz, 1003 Manila • Trunk Line 651-7800 local 1108, 1111, 1112, 1113
Direct Line: 711-9502; 711-9503 Fax: 743-1829 • URL: http://www.doh.gov.ph; e-mail: ftduque@doh.gov.ph

PS / COBAC
APPROVED PO / CONTRACT
Received by: _____
Date: AUG 07 2018

PS / COBAC
RELEASED: PO / CONTRACT
by: _____
Date: AUG 13 2018