

## Certificate of Good Manufacturing Practices


This certificate conforms to the format recommended by the World Health Organization

- 1. Purpose of this certificate:** To be introduced to Ministry of Health of Turkmenistan for the purpose of Export.
- 2. Certificate No.:** 1218/2021
- 3. Name and address of site:** Pharco B International (Parts no. 10,11,12,13,19,20 – block no. 18 - 3<sup>rd</sup> Industrial zone - New Borg El Arab City - Alexandria).
  - On the basis of the inspection, we certify that the site indicated on this certificate complies with Good Manufacturing Practices for the dosage forms, categories and activities listed in the Table 1 below
- 4. Manufacturer's license number:** 118042702008519 (License issued from Industrial Development Authority)

Dosage form(s)	Last Inspection	Category(ies)	Activities carried out by the company
<b>Non sterile:</b> Cephalosporin Solid dosage forms include: (Tablets – Hard gelatin capsules – dry syrup) <b>Sterile:</b> Filling of Cephalosporin vials	2019	General Category(ies)	Production and packaging
<b>Sterile:</b> (water for injection – ampoules – large volume parenteral “L.V.P” – Filling of Penem vials	2021		

- The responsibility for the quality of the individual batches of the pharmaceutical products manufactured through this process lies with the manufacturer.
- This certificate remains valid until 6/12/2022 It becomes invalid if the activities and/or categories certified herewith are changed or if the site is no longer considered to be in compliance with GMP.

Dr. Amira Mahgoub

  
Manager of the Inspection Administration of  
Human, Herbal and Veterinary and Disinfectants  
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Authenticated  
Dr. Yasin Ragaey

  
Head of Central Administration of Operations

Note: Not valid without stamp