


## 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 **United Arab Emirates** for the purpose of Registration.
- 2. Certificate No.:** 8/2022
- 3. Name and address of site** Egyphar (El Obour City, Industrial Zone(A) Block 13026, Parts No. 9 &10).
- On the basis of the inspection 2019 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:** 4 repeated/2007

Dosage form(s)	Category(ies)	Activities carried out by the company
<b>Non-sterile: Solid dosage form include :( Tablets – Hard gelatin Capsules - Powders).</b> <b>Liquid dosage form include: (Syrup).</b> <b>Soft gelatin capsule</b>	<b>General Category(ies)</b>	<b>Production and packaging</b>

- 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 13/1/2023 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  
Pharmaceuticals Factories.

Authenticated  
Dr. Yasin Ragaey  
  
Head of Central Administration of Operations

Note: Not valid without stamp