


Certificate of Good Manufacturing Practices

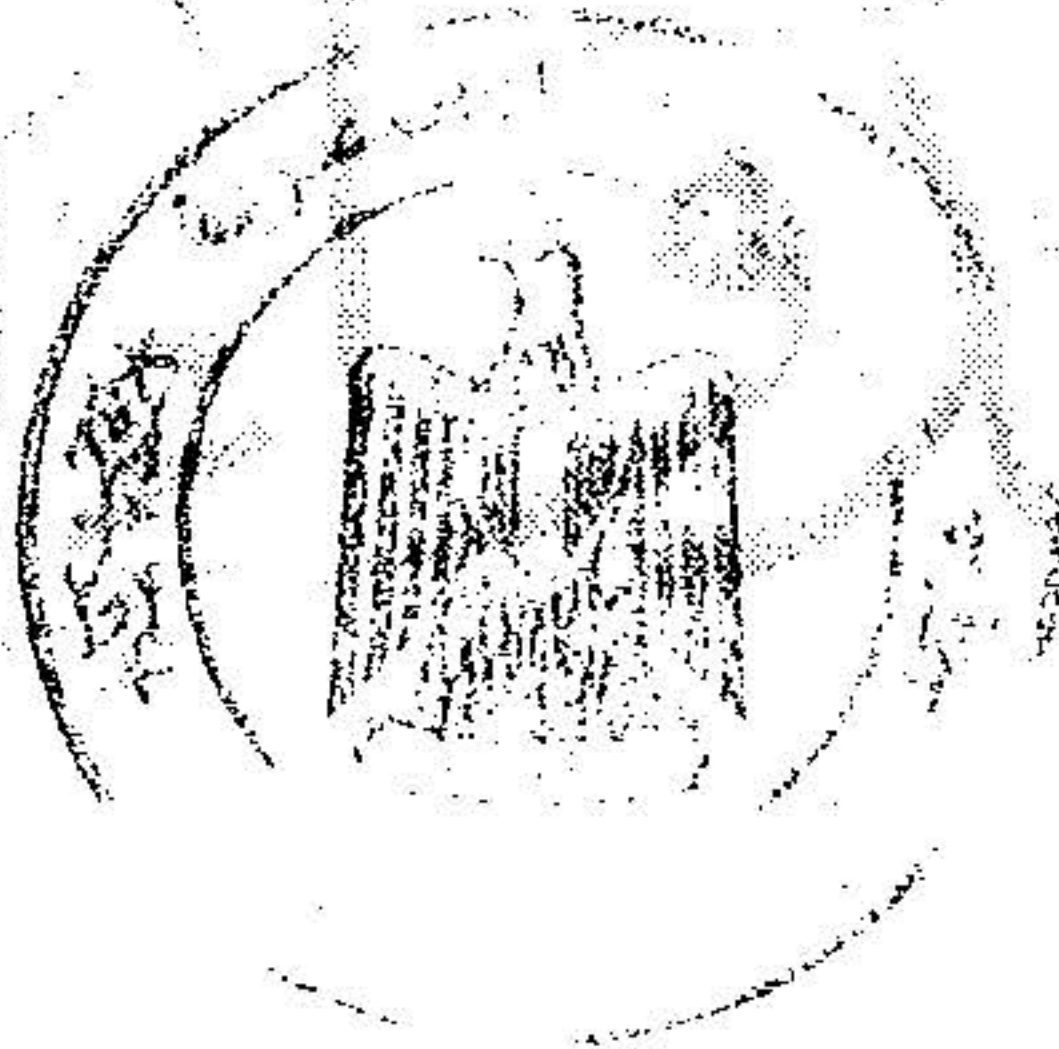
This certificate conforms to the format recommended by the World Health Organization

- 1. Purpose of this certificate:** To be introduced to Ministry of Health of **Morocco** for the purpose of Export.
- 2. Certificate No.:** 1237/2021
- 3. Name and address of site:** Chemipharm Pharmaceutical Industries (part no. 3/1 ,6th Industrial zone, 6th of October city – Giza).
• 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:** 112019033100008 (License issued from Industrial Development Authority)

Dosage form(s)	Last inspection	Category(ies)	Activities carried out by the company
<u>Sterile:</u> Cephalosporin (vial)	2019	General Category(ies)	Production and packaging
<u>Non-sterile:</u> Cephalosporin: Solid dosage forms ((Tablet – Hard Gelatin Capsules – powder for oral Suspension)			
<u>Non-sterile:</u> Solid dosage forms (Tablets – Hard Gelatin Capsules). <u>Liquid</u> (Syrup – Solution – Suspension). <u>Semisolid</u> (Ointment – Creams). <u>Sterile:</u> Ampoules & vials (solutions - lyophilized powder) - Eye drops	7/2021		

- 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
Pharmaceuticals Factories.



Authenticated
Dr. Yasin Ragaey

Head of Central Administration of Operations 

Note: Not valid without stamp