

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 Saudi Food and Drug Administration of Kingdom of Saudi Arabia for the purpose of Export
- 2. Certificate No.:** 1143/2021
- 3. Name and address of site:** Horis for Pharmaceutical Industries 2 Part no. 27 north extension area – Polaris international industrial parks – 6th of October city.
 - On the basis of the inspection 2021 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:** 414217
- 5. Table 1:**

Dosage Form(s)	Category(ies)	Activities carried out by the company
Non sterile, Solid dosage forms (Tablets – Hard Gelatin Capsule) - Powder filled in Sachet - Sterile Product (Sterile area includes Cephalosporin vial powder)	General Category(ies)	Production and packaging

- 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 16/11/2022 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 inspection Administration
of pharmaceutical, herbal and veterinary
preparations and disinfectants Factories



Authenticated:

Dr. Yasin Ragaey

Head of Central Administration of Operations

Note: Not valid without stamp

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