

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 Yemen for the purpose of Registration & Export.
- 2. Certificate No.:** 1346/2021
- 3. Name and address of site:** Badr pharma for Pharmaceutical Industries (Parts no. 164.165.166.167-zone of 250 feddans - Industrial zone – Badr city) .
- 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:** 1218042702011069 (License issued from Industrial Development Authority)

Dosage form(s)	Category(ies)	Activities carried out by the company
<u>Non-Sterile: Solid dosage forms</u> (Hard Gelatin Capsules –Dry mix in bottles) –Veterinary powder filled in sachets. <u>Liquid dosage forms</u> (Syrup – suspension – emulsion). <u>Cephalosporin area:</u> Solid dosage forms (powder for oral suspension). <u>Sterile: Non-β Lactam area</u> (Liquid in vial) - <u>Cephalosporin area:</u> vial (powder)	General Category(ies)	Manufacture 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 30/12/2022 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