

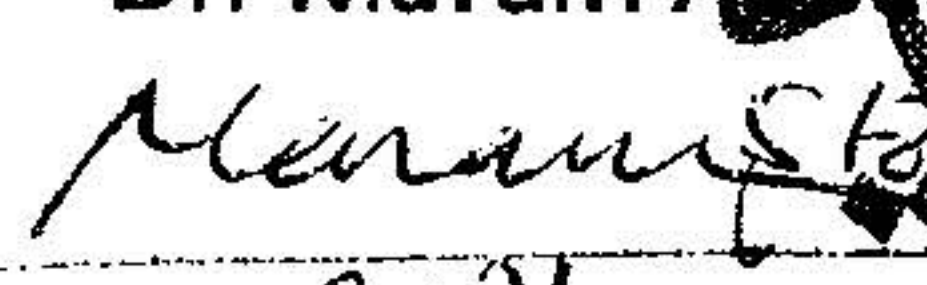
Certificate of Good Manufacturing Practices

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 **Libya** for the purpose of Registration & Export.
- 2. Certificate No.:** 846/2021
- 3. Name and address of site:** Badr pharma for Pharmaceutical Industries (Parts no. 164.165.166.167-industrial zone – Badr city) .
 - On the basis of the inspection carried out on 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:** 74/2019
- 5. Table 1:**

Dosage form(s)	Category(ies)	Activities carried out by the company
<u>Non-Sterile:</u> Solid (hard gelatin capsules –dry mix in bottles) – <u>Veterinary:</u> powder filled in sachets – Liquid (Syrup – suspension – emulsion) cephalosporin area: Solid (powder for oral suspension) – <u>Sterile:</u> Non-β lactam area (Liquid in vial)- cephalosporin area (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 3/8/2022. 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. Maram Abbas Sayed

Deputy General Manager for
Pharmaceutical Factory inspection



Dr. Yasin Ragaey


• Head of Central Administration of Operations

Note: Not valid without stamp

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