



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 Export.
- 2. Certificate No.:** 452/2021
- 3. Name and address of site:** Organo for pharmaceutical & Chemical Industries (Organo Pharma) (Industrial zone - block 20005 - Part no. 2011 - El Obour city).
 - On the basis of the inspection carried out on 2020 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:** 202007
- 5. Table 1:**

Dosage form(s)	Category(ies)	Activities carried out by the company
Solid dosage forms (Tablet - Hard gelatin capsule) - Powder (Human & veterinary) - Liquid (Syrup - Suspension)	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 25/4/2022 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

Maram
2021
Deputy General Manager for
Pharmaceutical Factory inspection



Dr. Yasin Ragaey

Yasin Ragaey
27/4

Head of Central Administration of Operations

Note: Not valid without stamp

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