

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 CA of Biological and Innovative Products and Clinical Studies for Registration.
- 2. Certificate No.:** 865 /2021
- 3. Name and address of site:** Chemical Industries Development (CID) (El Aharam st – El talbyaa – El Giza).
 - 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:** 6 /1950
- 5. Table 1:**

| Dosage form(s) | Category(ies) | Activities carried out by the company |
|--|-----------------------|---------------------------------------|
| Sterile Products: Human ampoule (Terminal sterilization) | General Category(ies) | Manufacturing - 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 16/8/2022 and 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.

Handwritten signature and date: 7/9/2021

Dr. Omnia Medhat
Biological Inspection Lead
Handwritten signature and date: 7/9/2021

هيئة الدواء المصرية

Dr. Yasin Ragaey



Head of Central Administration of Operations

Note: Not valid without stamp

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