

## 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 Indonesia for the purpose of Registration & Export.

2. **Certificate No.:** 831/2021

3. **Name and address of site:** Liptis for pharmaceuticals and Medical products (plot No. B/2-6<sup>th</sup> industrial zone-4<sup>th</sup> of October City-Giza).

• On the basis of the inspection carried out on 5/2019 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:** 1-20180-280-018 (License issued from Industrial Development Authority)

5. **Table 1:**

Dosage form(s)	Category(ies)	Activities carried out by the company
Human: <u>Non-Sterile</u>	General	Production and Packaging
Solid (Tablets filled in blisters & jars – Hard Gelatin Capsules filled in blisters & jars	Category(ies)	

• 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 2/8/2022 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.

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*Maram*  
Deputy General Manager for  
Pharmaceutical Factory Inspection

هيئة الدواء المصرية  
*Yasin Ragaey*  
2/8

Dr. Yasin Ragaey

Head of Central Administration of Operations

Note: Not valid without stamp

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