

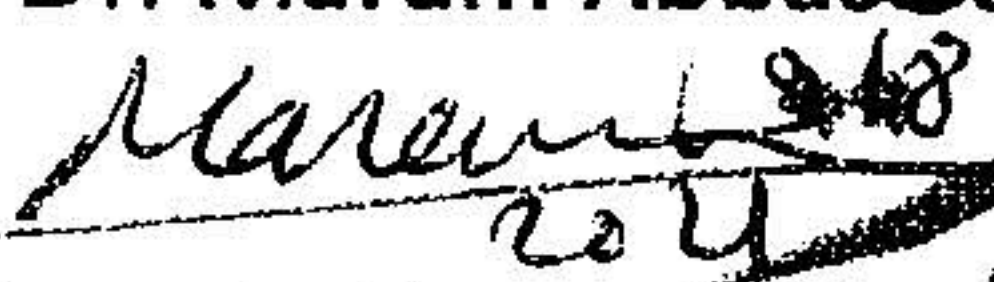
Certificate of Good Manufacturing Practices


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 Nigeria for the purpose of Export.
- 2. Certificate No.:** 850/2021
- 3. Name and address of site:** Accidima International Trading (Plot no. 41-6th Industrial zone extension- 6th October 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:** 1-20190-210-002 (License issued from Industrial Development Authority)
- 5. Table 1:**

Dosage form	Category(ies)	Activities carried out by the company
Compressed Aerosol (Powder-Liquid - foam) – Non-Compressed Aerosols (Lotion - Liquid) – Solid Dosage forms include Hormonal Tablet (coated – uncoated – sugar & film coat") –Sterile: Hormonal ampoule (liquid & suspension) and Hormonal Vial (liquid – suspension)	General Category(ies)	Manufacturing 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 5/8/2022. 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

هيئة الدواء المصرية

Head of Central Administration of Operations

Note: Not valid without stamp

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