

## 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 Kenya for the purpose of Registration & Importation.
- 2. Certificate No.:** 929/2021
- 3. Name and address of site:** Sanofi Egypt (No. 3 El Massaneh st. El Mirya – El Zeitoun – Cairo – Egypt).
  - On the basis of the inspection 2/2019 we certify that the site indicated in 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:** 2180/7020/145 (License issued from Industrial Development Authority)

Dosage form(s)	Category	Activities carried out by the company
<u>Non-sterile:</u> Solid (Tablet – Capsule)- Liquid (Syrup – Suspension) Semi-solid (Ointment – Cream – Gel)- Suppositories – <u>Sterile:</u> Cephalosporin Amp (vial) Ampoule (Terminal sterilized)	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 20/3/2022 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. Amr Hagazi

Factories inspection General Manager

Authenticated:

Dr. Yasin Ragaey

Head of Central Administration of Operations

Note: Not valid without stamp

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