Arab Republic of Egypt
Egyptian Drug Authority
Central Administration of Operations
Factories of pharmaceutical, herbal
And veterinary preparations and
disinfectants

جمهورية مصر العربية
هيئـة الدواء المصرية
الإدارة المركزية للعمليات
مصانع المستحضرات الصيدلية والعشبية
والبيطرية والمطهرات

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 **France** for the purpose of Registration & Import.
2. **Certificate No.:** 689/2021
3. **Name and address of site:** Sanofi Egypt (No. 3 El Massaneh st. El Amirya – El Zeitoun – Cairo – Egypt).
   - **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. **Manufacturing license number:** 2180/2020/145 (License issued from Industrial Development Authority)
5. **Table 1:**

Dosage form(s)	Category(ies)	Activities carried out by the company
Non-sterile: Solid (Tablets – Capsule)- Liquid (Syrup – suspension) Semi-solid (Ointment – Cream – Gel)- Suppositories – Sterile: Cephalosporin Area (vial) -Ampoule (Terminally 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** 20/6/2022 **It becomes invalid if the activities and/or categories certified herewith are changed or if the site is no longer considered to be incompliance with GMP.**

Dr. Maram Abbas Sayed
Deputy General Manager for
Pharmaceutical Factory inspection

Dr. Yasin Ragaey
Head of Central Administration of Operations
22/6

Note: Not valid without stamp

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