

Certificate of Good Manufacturing Practices

This certificate conforms to the format recommended by the World Health Organization

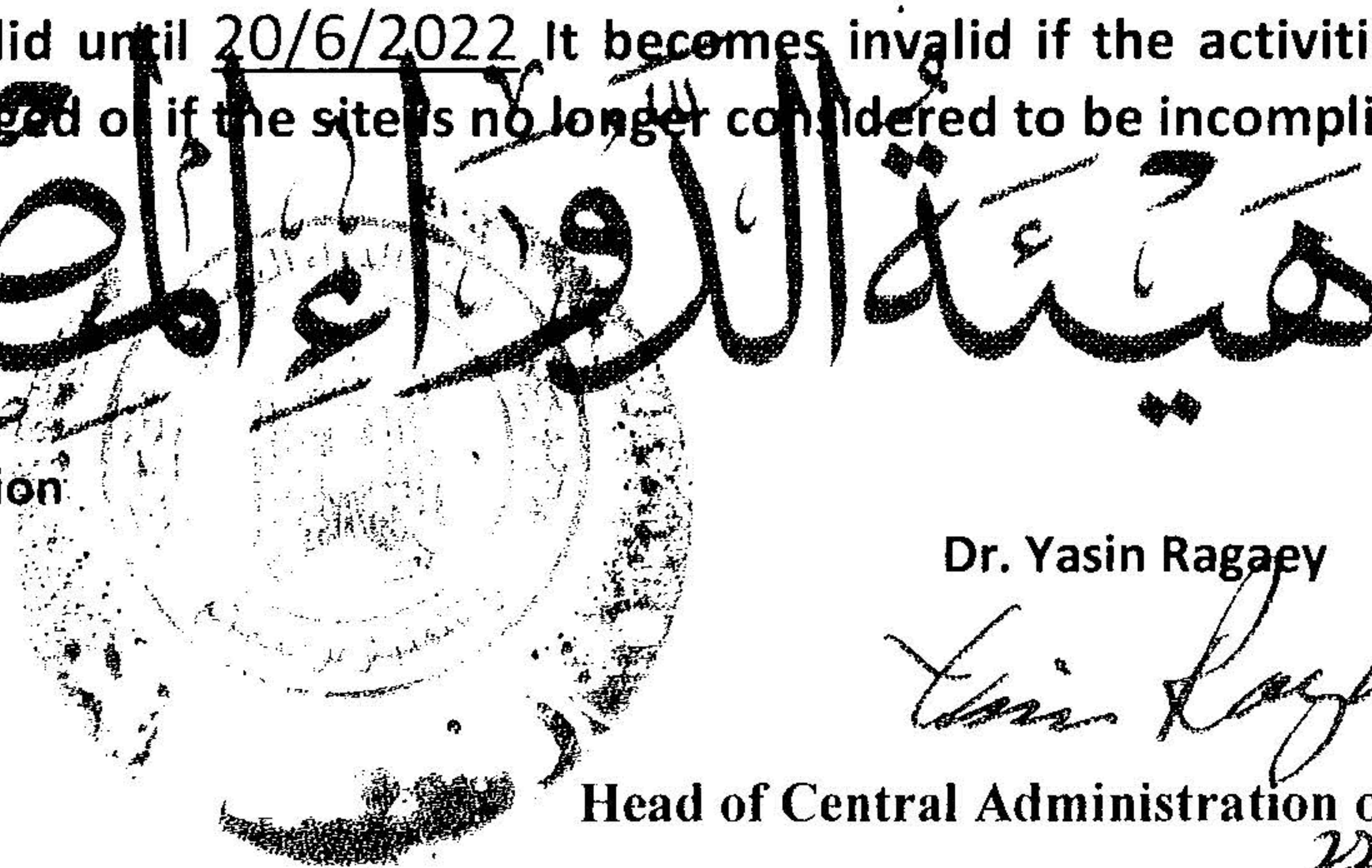
- 1. Purpose of this certificate:** To be introduced to Ministry of Health of France for the purpose of Registration & Import.
- 2. Certificate No.:** 689/2021
- 3. Name and address of site:** Sanofi Egypt (No. 3 El Massaneh st. El Mirya – El Zeitoun – Cairo – Egypt).
 - On the basis of the inspection 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. Manufacturer license number:** 2180-7020-143 (License issued from Industrial Development Authority)
- 5. Table 1:**

Dosage form (s)	Category(ies)	Activities carried out by the company
<u>Non-sterile:</u> Solid (Tablets – Capsule)- Liquid (Syrup – Suspension) Semi-solid (Ointment – Cream – Gel- Suppositories – <u>Sterile:</u> Cephalosporin Intra (vial) - Ampoules (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 here with are changed or if the site is no longer considered to be in compliance with GMP.

Dr. Maram Abbas Sayed
Maram
2021
Deputy General Manager for
Pharmaceutical Factory inspection

هيئة الدواء المصرية



Dr. Yasin Ragaey

Yasin Ragaey
Head of Central Administration of Operations

Note: Not valid without stamp

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