Arab Republic of Egypt
Egyptian Drug Authority
Central Administration of Operations
Biological Factories Inspection department

جمهورية مصر العربية
هيئـــة الـدواء المصرية
الإدارة المركزية للعمليات
ادارة التفتيش على مصانع المستحضرات الحيوية

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 **Palestine** for the purpose of Registration & Export.

2. **Certificate No.:** 1343/2021

3. **Name and address of site:** Wadi El Neel Benta for Pharmaceuticals and Medical Devices (Part no. 64-industrial zone 5A-10th of Ramadan City – El Sharkya).

- **On the basis of the inspection** 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:** 918042702008751(License issued from Industrial Development Authority)

5. **Table 1:**

Dosage form(s)	Category(ies)	Activities carried out by the company
Sterile: Prefilled Syringe	General Category(ies)	Manufacturing, Packaging and marketing

- **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** 28/12/2022 **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. Omnia Medhat

Dr. Omnia 28/12/2021
Biological Inspection Head

Authenticated:
Dr. Yasin Ragaey
Yasin Ragaey
Head of Central Administration of Operations
30/12

Note: Not valid without stamp

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