


Certificate of Good Manufacturing Practices

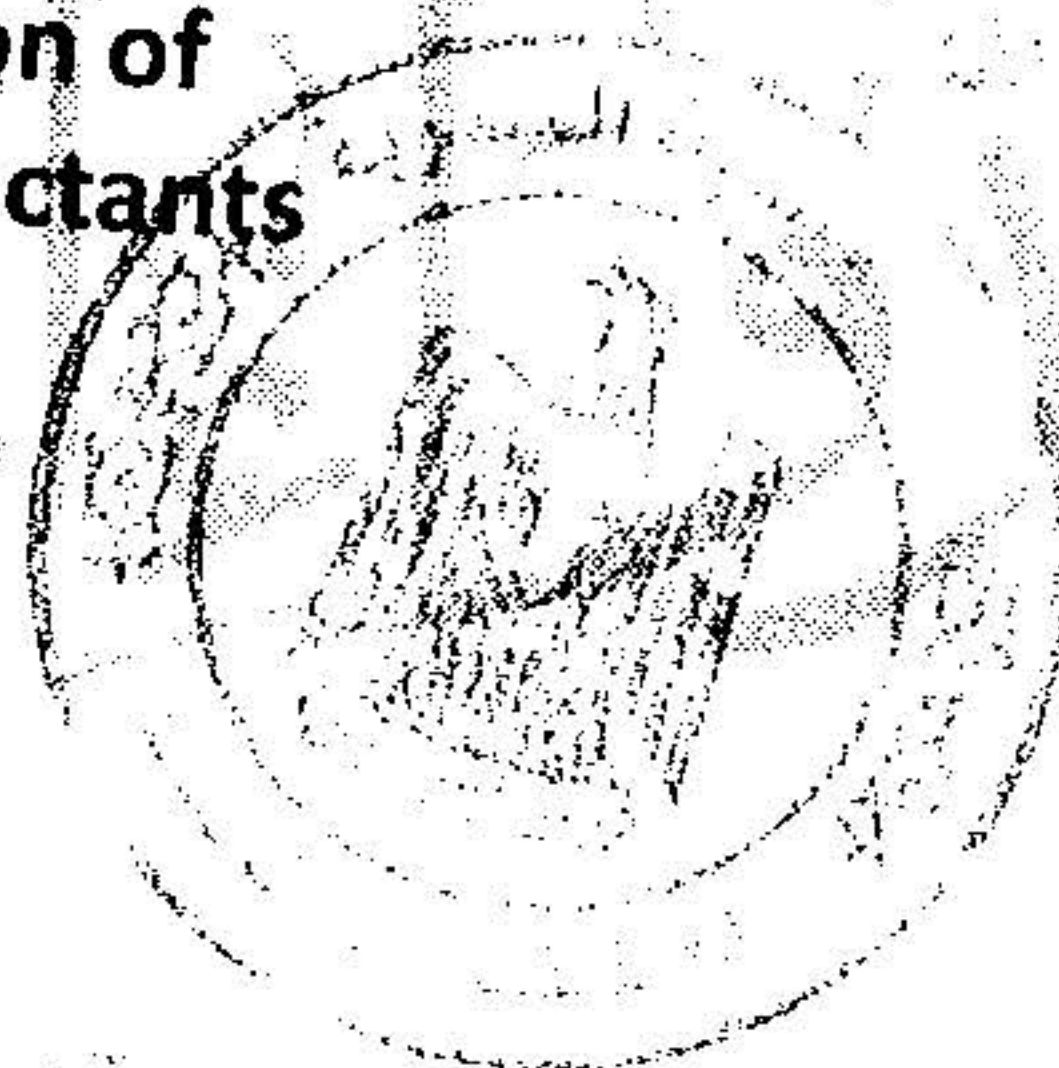
This certificate conforms to the format recommended by the World Health Organization

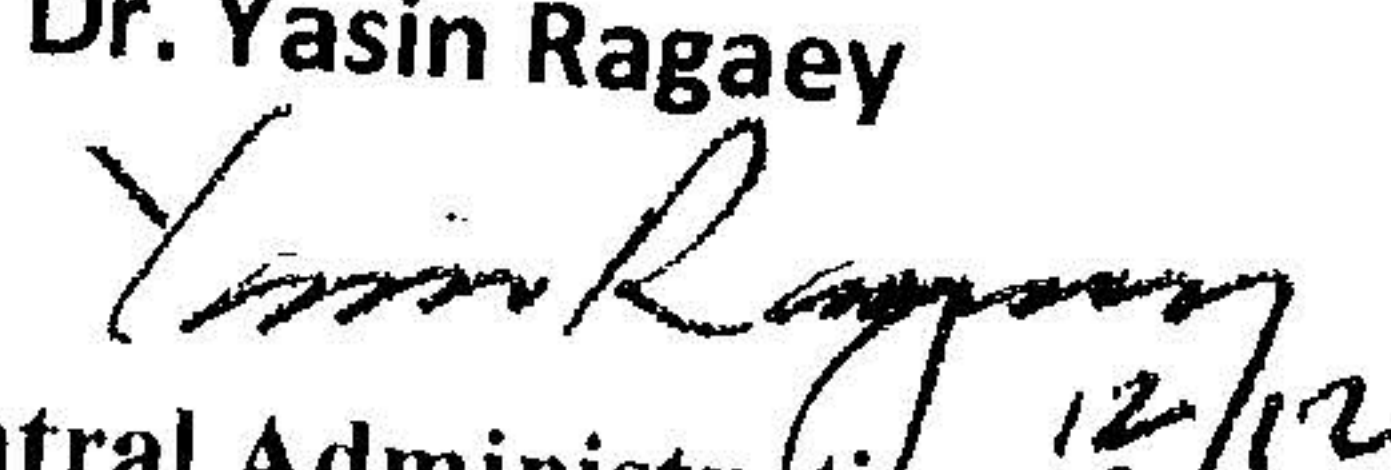
- 1. Purpose of this certificate:** To be introduced to Ministry of Health of Chile for the purpose of Export.
- 2. Certificate No.:** 1259/2021
- 3. Name and address of site:** Novartis Pharma S.A.E (No. 3 - El-Sawah st. - El-Amirya - Cairo).
• On the basis of the inspection, we certify that the site 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:** 134/1965
- 5. Table 1:**

Dosage form(s)	Last inspection	Category(ies)	Activities carried out by the company
Non-sterile: Solid dosage form (Tablet - Hard gelatin capsules - powder filled in sachets). Semisolid dosage forms (Suppositories).	2019	General	Production and packaging
Sterile: (Ampoule)	2021	Category(ies)	

- 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 9/12/2022 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. Amira Mahgoub

Manager of the Inspection Administration of
Human, Herbal and Veterinary and Disinfectants
Pharmaceuticals Factories.



Authenticated
Dr. Yasin Ragaey

Head of Central Administration of Operations

Note: Not valid without stamp