

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 Kazakhstan for the purpose of Registration & Export.
- 2. Certificate No.:** 1261/2021
- 3. Name and address of site:** Marcyrl Pharmaceuticals Industries (MPI) (Parts no. 5,6,7,8,9,10,11,12 - west Extension of the industrial zone, Block 20005 - El Obour City).
- On the basis of the inspection 2020 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:** 112019021700002 (License issued from Industrial Development Authority)

5. Table 1:

Dosage form(s)	Category(ies)	Activities carried out by the company
<u>Non-Sterile: Human: Solid dosage forms (Tablets – Hard Gelatin Capsules - Powders Packed in sachets).</u> <u>Liquids dosage forms (Syrup – Oral Drops).</u> <u>Semisolid dosage forms (Creams – Ointments – Gel).</u> <u>Non hormonal Suppositories - Soft Gelatin capsules.</u> <u>Solid dosage forms (Tablets– Bilayer Tablets)</u>	General Category(ies)	Manufacture 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 13/12/2022 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
Amira Mahgoub
Manager of the Inspection Administration of
Human, Herbal and Veterinary and Disinfectants
Pharmaceuticals Factories.

Authenticated
Dr. Yasin Ragaey
Yasin Ragaey
Head of Central Administration of Operations
13/12

Note: Not valid without stamp