

## Certificate of Good Manufacturing Practices

This certificate is issued as per WHO TRS No 908 of 2003

- 1. Purpose of this certificate:** To be introduced to Ministry of Health of Iraq for the purpose of Export.
- 2. Certificate No:** 287/2022
- 3. Name and address of site:** Euromed for medical industries –free zone,-Nasr city, part no.10-block (I).  
P.O. BOX:11816, Cairo, Egypt
- 4. Manufacturer's license number:** 21/2002 (expired on 16/7/2027)
- 5. On the basis of the inspection on 17/06/2021 we certify that the site indicated on this certificate complies with Good Manufacturing Practices for the activities listed in the Table 1 below and list of products described in products list attached to this Certificate**

**Table 1:**

Production line	Activities
Syringe	Injection, printing, Assembly, Packaging, packing Sterilization
Cannula	Imported component, Injection, Assembly, Packaging, packing - Sterilization
Packaging of Catheter	Imported non sterile devices, packaging, packing - Sterilization
Packaging of tracheal tubes (with cuff- without cuff)	Imported non sterile devices, packaging, packing - Sterilization

- The responsibility for the quality of the individual batches of the medical devices manufactured through this process lies with the manufacturer.
- This certificate remains valid until 15/06/2022 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. Sally Abd El Rasoul

Medical Devices & Invitro diagnostic inspection manager

Authenticated:  
Dr. Yasin Ragaey

Head of Central Administration of Operations

Note: Not valid without stamp

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