

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 UP-MS-MT for dialysis tender.
- 2. Certificate No:** 1180/2021
- 3. Name and address of site:** Afri Medical-industrial area c3-10th of Ramadan city
- 4. Manufacturer's license number:** 609
- 5. On the basis of the inspection 29/11/2021 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 with this Certificate**

Table 1:

Production line	Activities
Hemodialysis & its substitutes	manufacturing - packaging - 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 28/09/2022. 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
Sally Abdel Rasoul
24/11/2021
Medical Devices & Invitro diagnostic inspection manager

Authenticated:
Dr. Yasin Ragaey

Head of Central Administration of Operations

هيئة الدواء المصرية

Note: Not valid without stamp

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