

LETTER OF AUTHORIZATION

To whom it may concern,

We, **KENMAK HASTANE MALZEMELERİ ELEKTROSTATİK BOYA SAN. TİC. A. Ş**
on business address (Dokuz Eylül, Sarnıç Yolu No:23, 35410 Gaziemir/İzmir, herewith
authorize

M-INTER-FARMA S.A. on business address str. Grenoble 23. mun. Chişinau MD2025,
Republic of Moldova

- As our **Authorized Representative** (in correspondence with the conditions of directive 93/42/EEC) for Notification/Registration of medical devices to the Agency of the Medical Products and Medical Devices and the inclusion in the Register of the medical devices manufactured by **KENMAK HASTANE MALZEMELERİ ELEKTROSTATİK BOYA SAN. TİC. A. Ş**, in territory of Republic of Moldova.
- We declare that the company mentioned above is authorized to register, notify, renew or modify the registration of medical devices on the territory of the Republic of Moldova, and to perform Essential Duties required by Law No. 102 09.06.2017 regarding medical devices.
- as our Authorized Representative to promote, advertise, participate in public tenders and subsequently negotiate and sign Contracts, to make delivery, installation, training of the personnel for work with the equipment and to perform warranty and after warranty service for the products of our manufacture on the territory of Moldova.

As the manufacturer, we guarantee our products against defects in materials and workmanship, and provide services based on the standard terms and conditions of our warranty policy.

This Authorization letter is valid until 31th December, 2025 or till conclusion of all contracts for our products signed in 2023-2025.

Date 08 december 2023

stamp

Signature

Signature and stamp

Mr. YAMAN BAŞ

CEO –


KENMAK
HASTANE MALZEMELERİ ELEKTROSTATİK
BOYA SANAYİ TİCARET LİMİTED ŞİRKETİ
Dokuz Eylül Mh. Sarnıç Yolu No:23 Gaziemir - İZMİR
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Mersis No: 354052745900001
Gaziemir V.D. 544 062 7459

Vers 01.01.2023