

CERTIFICATE

The Certification Body TÜV Rheinland Italia S.r.l.

certifies, in accordance with the TÜV Rheinland Group procedures, that the Company

DISPOTECH S.R.L.

Via al Piano, 29

IT – 23020 Gordona (SO)



has established and applies a quality management system for the following scope:

Manufacturing and placing on the market of medical devices in laminated material tissue/polyethylene and devices for thermal therapy. Management of manufacture and placing on the market of ice spray and saliva ejector.

Through an Audit, Report No. 7981994130SAN27, proof has been furnished that the quality management system fulfils the requirements of the standard

UNI CEI EN ISO 13485:2021

Please refer to the Quality Manual for the details about the exclusions with respect to the requirements of the standard.

Certificate Registration No. **39 05 1040609**

This Certificate is valid from 2025/01/28 to 2028/01/27

The reference date for all the next audits is (day-month): 14/01

Milan, 2025/01/24. First Certification: 2004/12/15

The certification responsible: Cesare Gentile
TÜV Rheinland Italia S.r.l., Via E. Mattei, 3 - I - 20005 Pogliano Milanese (MI)

This certificate does not represent proof that the statutory requirements of the Directives 93/42/EEC, 90/385/EEC, 98/79/EC or Regulations (UE) 2017/745, (UE) 2017/746 have been fulfilled



MS N° 0083

Membro degli Accordi di Mutuo Riconoscimento EA, IAF e ILAC.
Signatory of EA, IAF and ILAC Mutual Recognition Agreements.



Management System
EN ISO
13485:2016

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