

CERTIFICATE

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

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

Codan S.r.l.

Via XXV Aprile, 161

IT – 41037 Mirandola (MO)

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

Distribution of medical devices active and non-active sterile and not sterile for infusion, injection and transfusion. Distribution of active medical devices sterile and not sterile for the monitoring of vital physiological parameters. Import of active medical devices not sterile for infusion therapy. Provision of maintenance service for infusion.

Through an Audit, Report No. 7994166010FES06, 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 0401903**

This Certificate is valid from 2025/05/14 to 2028/05/13

The reference date for all the next audits is (day-month): 29/03

Milan, 2025/05/09. First Certification: 2019/05/14



The certification responsible: Daniele Ricchi
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
+AC:2018+A11:2021
ISO 9001:2015

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