



America

CERTIFICATE

No. QS6 047541 0020 Rev. 03

Certificate Holder:

Cedic s.r.l.
Via Liberazione 63/ 9
20068 Peschiera Borromeo (MI)
ITALY

Certification Mark:



Scope of Certificate:

Design and Development, Production and Distribution of Sterile Disposable Catheters, Sets for Pump or Gravity, Extension Sets, Connectors, Adapters and Bags for Enteral Nutrition, Enteral Drug Delivery and Gastric Residuals Drainage

Standard(s):

ISO 13485:2016

Regulatory Authority(ies):

Australia TGA, Brazil ANVISA, Health Canada, Japan MHLW / PMDA, USA FDA. See attached for listing of specific regulatory requirements.

The Certification Body of TÜV SÜD America Inc. certifies that the quality management system of the manufacturer listed above has been audited against the stated criteria and found to conform to those criteria for the scope of certification listed. For details and certificate validity see:

www.tuvsud.com/ps-cert?q=cert:QS6 047541 0020 Rev. 03

TÜV SÜD America Inc. is an MDSAP Recognized Auditing Organization.

REPs Facility ID:

F001656

Report No.:

ITA200220005497

Effective Date:

2025-04-07

Expiry Date:

2027-04-07

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(Renee Walker)
Director, US Certification Body, MHS

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Regulatory Requirements: Audit/Certification Criteria

Australia

Therapeutic Goods (Medical Devices) Regulations 2002
- Schedule 3, Part 1 (excluding Part 1.6) – Full Quality Assurance Procedure

Brazil

- RDC ANVISA n. 665/2022 - Good Manufacturing Practices
- RDC ANVISA n. 551/2021
- RDC ANVISA n. 67/2009 - Vigilance

Canada

- Medical Device Regulations – Part 1- SOR 98/282

Japan

- MHLW Ministerial Ordinance No. 169 (2004), as amended by MHLW Ministerial Ordinance No.60 (2021)
- Japan PMD Act (as applicable)

United States

- 21 CFR Part 803
- 21 CFR Part 806
- 21 CFR Part 807 – Subparts A to D
- 21 CFR Part 820

Facility(ies):

Cedic s.r.l.

Via Liberazione 63/ 9, 20068 Peschiera Borromeo (MI), ITALY

Facility Scopes:

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