

CERTIFICATE

Number: 2229417

The management system of:

Technomed Europe

Wiebachstraat 25a
6466 NG Kerkrade
Netherlands

Manufacturer Facility Identifier F002732

Conforms with the following standard and regulatory requirements:

ISO 13485:2016

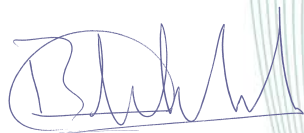
Australia: Therapeutic Goods (Medical Devices) Regulations, 2002 and Schedule 3 Part 1 (excluding Part 1.6) - Full Quality Assurance Procedure
Brazil: RDC ANVISA n. 665/2022, 551/2021 and 67/2009
Canada: Medical Devices Regulations - Part 1 - SOR 98/282
Japan: MHLW Ministerial Ordinance 169, Article 4 to Article 68 and PMD Act
United States: 21 CFR 803, 21 CFR 806, 21 CFR 807 - Subparts A to D and 21 CFR 820

Scope:

The design and development, manufacture, and distribution of sterile and non-sterile stimulating and recording electrodes, including EEG electrodes, adhesive electrodes, needle electrodes, probes, surgical instruments, and laryngeal electrodes for neuro-diagnostic examinations; including extension cables; and the distribution of neuro-diagnostic systems for intra-operative monitoring

Certificate expiry date: 2029-05-01
Certificate effective date: 2026-05-01
Certified since: 2020-05-14

DEKRA Certification B.V.



B.T.M. Holtus
Managing Director



F. Godeke
Certification Manager

© Integral publication of this certificate and adjoining reports is allowed.

The validity of this certificate can be checked through DEKRA's website using the following link:
<https://www.dekra-checkme.com/org>

DEKRA Certification B.V. is recognized under the Medical Devices Single Audit Program.



DEKRA Certification B.V. Meander 1051, 6825 MJ Arnhem P.O. Box 5185, 6802 ED Arnhem, Netherlands
T +31 88 96 83000 F +31 88 96 83100 www.dekra.nl Company registration 09085396