EC CERTIFICATE

Number: 2109122CE01

Full Quality Assurance System

Directive 93/42/EEC on Medical devices, Annex II excluding (4)
(Devices in Class IIa, IIb or III and Devices in Class I in sterile conditions and sterilised systems or procedure packs)

Manufacturer:

Technomed Europe

Amerikalaan 71
6199 AE Maastricht Airport
The Netherlands

For the product category(ies)

Stimulation-, needle-, gel electrodes and accessories to the applicable equipment including sterile extension cables

DEKRA grants the right to use the EC Notified Body Identification Number illustrated below to accompany the CE Marking of Conformity on the products concerned conforming to the required Technical Documentation and meeting the provisions of the EC-Directive which apply to them:

0344

Documents, that form the basis of this certificate:

Certification Notice 2109122CN, initially dated 28 November 2007
Addendum, initially dated 28 November 2007

DEKRA hereby declares that the above mentioned manufacturer fulfils the relevant provisions of 'Besluit Medische Hulpmiddelen', the Dutch transposition of the Council Directive 93/42/EEC of June 14, 1993 concerning Medical devices, including all subsequent amendments. The manufacturer has implemented a quality assurance system for design, manufacture and final inspection, that covers the aspects of manufacture concerned with securing and maintaining sterile conditions, for the above mentioned product category in accordance to the provisions of Annex II Council Directive 93/42/EEC of June 14, 1993 and is subject to periodical surveillance. For placing on the market of Class III devices an additional EC design examination certificate according to Annex II (4) is mandatory.
The necessary information related to the quality management system of the manufacturer, including facilities and the reference to the relevant documentation, of the products concerned and the assessments performed, are stated in the Certification Notice which forms an integrative part of this certificate.

This certificate is valid until: 1 November 2023
Issued for the first time: 28 November 2007
Revised: 21 January 2015
Reissued: 20 November 2019

DEKRA Certification B.V.

B.T.M. Holtus
Managing Director

J.A. van Vugt
Certification Manager

DEKRA Certification B.V. is Notified Body with ID no 0344

DEKRA Certification B.V. Meander 1051, 6825 MJ Arnhem P.O. Box 5185, 6802 ED Arnhem, The Netherlands
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ADDENDUM

Belonging to certificate: 2109122CE01

CE MARKING OF CONFORMITY
MEDICAL DEVICES

Stimulation-, needle-, gel electrodes and accessories to the applicable equipment including sterile extension cables

Issued to:

Technomed Europe
Amerikalaan 71
6199 AE Maastricht Airport
The Netherlands

This certificate covers the following product(s):

Sterile disposable Bipolar and Monopolar stimulating electrodes and needle electrodes for neurophysiological examinations such as EEG, EMG procedures

Sterile disposable Bipolar and Monopolar stimulating electrodes for the location of various nerves during surgical procedures

Sterile extension cables for stimulation-, needles-, gel electrodes connected to the applicable equipment

Initial date: 28 November 2007
Revision date: 20 November 2019

DEKRA Certification B.V.

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