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EC Certificate

Full Quality Assurance System Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4) (Devices in Class IIa, IIb or III)

No. G1 089300 0006 Rev. 00

Manufacturer:

neuroConn GmbH

Albert-Einstein-Str. 3 98693 Ilmenau **GERMANY**

Facility(ies):

neuroConn GmbH

Albert-Einstein-Str. 3, 98693 Ilmenau, GERMANY

Product Category(ies): Stimulators for Cranial Electrotherapy, **Biofeedback Systems**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.:

713163181

Valid from:

2019-08-27

Valid until:

2024-05-26

Date,

2019-08-27

Stefan Preiß

1. Punil

Head of Certification/Notified Body