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Product Service

# EC Certificate

## Product Quality Assurance System

**Directive 93/42/EEC on Medical Devices (MDD), Annex VI**
**(Devices in class IIa or IIb)**

**No. G3 17 08 48742 013**

**Manufacturer:**

**MTR plus Vertriebs GmbH**
Kamenzer Damm 78
12249 Berlin
GERMANY

**Facility(ies):**

MTR plus Vertriebs GmbH
Kamenzer Damm 78, 12249 Berlin, GERMANY

**Product Category(ies):**

**Electrotherapy devices for nerve- and muscle-stimulation, EMG-Biofeedback units and vaginal- and rectal probes**

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

**Report no.:** 713115190

**Valid from:** 2017-12-20
**Valid until:** 2022-12-19

**Date,** 2017-11-14

Stefan Preiß

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123







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