



Product Service

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 11 03 31461 032

Manufacturer: BMT Medical Technology s.r.o

Cejl 50
656 60 Brno
CZECH REPUBLIC

Facility(ies):

BMT Medical Technology s.r.o
Cejl 50, 656 60 Brno, CZECH REPUBLIC

**Product
Category(ies):**

**Devices for Sterilization and
Disinfection in Health Care Facilities**

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.: 71382413

Valid from: 2011-03-29

Valid until: 2015-01-31

Date, 2011-04-08

Hans-Heiner Junker



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

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