ZERTIFIKAT ◆ CERTIFICATE ◆ 認証書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

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 14 10 22725 017

Manufacturer:

MMM Münchener Medizin Mechanik GmbH
Semmelweisstraße 6
82152 Planegg
GERMANY

Facility(ies):

MMM Münchener Medizin Mechanik GmbH
Semmelweisstraße 6, 82152 Planegg, GERMANY

MMM Münchener Medizin Mechanik GmbH
Hauptstraße 2, 92549 Stadlern, GERMANY

MMM Münchener Medizin Mechanik GmbH
Zechenstrassse 12, 86971 Peiting, GERMANY

Product Category(ies):

Devices and Systems for Sterilization, Disinfection and Processing 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.: 713051216

Valid from: 2015-01-21
Valid until: 2020-01-20

Date, 2014-12-17

Hans-Heiner Junker

595042

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

TÜV SÜD Product Service GmbH · Zertifizierstelle · Ridlerstraße 65 · 80339 München · Germany

TÜV®