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TÜV SÜD 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 15 04 53618 020

Manufacturer:	**ZHERMACK S.p.A** Via Bovazecchino, 100 45021 Badia Polesine (RO) ITALY
Facility(ies):	ZHERMACK S.p.A Via Bovazecchino, 100, 45021 Badia Polesine (RO), ITALY
Product Category(ies):	**Disinfectants for medical devices;** **Acrylic resins for dentures;** **Dental restorative materials;** **Steam Sterilizing unit**

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.: ITA258960

Valid from: 2015-05-06
Valid until: 2020-05-05

665034

Date, 2015-05-05

Hans-Heiner Junker

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

Zhermack S.p.A