

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 08 15409 023

Manufacturer: Ivoclar Vivadent GmbH
Dr. Adolf-Schneider-Str. 2
D-73479 Ellwangen
Deutschland
Facility(ies): Ivoclar Vivadent GmbH



Dr. Adolf-Schneider-Str. 2, D-73479 Ellwangen, Deutschland

Product
Category(ies): Dental instruments, Dental medical devices

The Certification Body of TUV SUD 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.: 713067667

Valid from: 2016-01-16
Valid until: 2021-01-15

Date, 2015-12-22


Hans-Heiner Junker

