Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

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 095377 0008 Rev. 00

Manufacturer:

Prevest Denpro Limited
Export Promotion Industrial Park
(EPIP), Bari Brahamana
181133 JAMMU
INDIA

Product Category(ies): Dental Materials (Class IIa):
Restorative Materials (Filling, core build up and luting materials) Sealants, Liners and Base Materials, Adhesives, Cements and Filling Materials, Zinc Oxide-Eugenol Temporary Cements, Non-Eugenol Temporary Cements, Root Canal Sealing Materials, Endodontic Materials, Pulp Capping Materials, Varnishes and Desensitizers, Denture Base Polymers, Etching Materials, Auxiliary Materials for Dental Treatment.

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.: IND2019072

Valid from: 2020-04-23
Valid until: 2024-05-26

Date, 2020-04-23

Christoph Dicks
Head of Certification/Notified Body

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

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany