



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