



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 16 06 95378 001

Manufacturer: Prevest Denpro Limited

38 Industrial Estate
Digiana
180010 Jammu
INDIA

EC-Representative: Obelis s.a

53, Boulevard General Wahis
1030 Brussels
BELGIUM

**Product
Category(ies):**

Dental Materials:
Restorative Materials (Filling, core build up
and luting materials)
Sealants, liners and base materials,
Adhesives, cements, Temporary
Non eugenol, Cements and filling materials,
root canal sealing materials,
Endodontic materials,
Pulp Capping materials,
Varnishes and desensitizers,
denture base polymers,
etching materials,
auxillary 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.

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Date, 2017-01-09

Stefan Preiß



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

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