



EC Certificate

Production Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex V
(Devices in Class IIa, IIb or III)

No. G2 090341 0009 Rev. 01

Manufacturer:

Candulor AG

Boulevard Lilienthal 8
8152 Glattpark (Opfikon)
SWITZERLAND

**Product
Category(ies):**

**Acrylic and ceramic teeth,
denture base materials for fixed dentures,
denture base colorants for fixed dentures**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex V. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class IIb and III devices an additional Annex III certificate is mandatory. All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:G2_090341_0009_Rev.01

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Valid until: 2024-05-26

Date, 2021-03-18

Christoph Dicks
Head of Certification/Notified Body