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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 17 09 93599 005

**Manufacturer:**

**C-TECH IMPLANT SRL**
Via Cesare Battisti 2
40123 Bologna
ITALY

**Facility(ies):**

C-TECH IMPLANT S.r.l.
Via San Benedetto 1837, 40018 San Pietro in Casale (BO), ITALY

**Product Category(ies):**

**Dental implants, accessories for dental implants**

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.:** ITA2771461S

**Valid from:** 2017-11-10
**Valid until:** 2020-11-08


**Date,** 2017-11-10

Stefan Preiß

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





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