



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 03 76870 011

Manufacturer: **ALTATEC GmbH**

Maybachstr. 5
71299 Wimsheim
GERMANY

Facility(ies):

ALTATEC GmbH
Maybachstr. 5, 71299 Wimsheim, GERMANY

ALTATEC GmbH
Paul Ehrlich Str. 15, 72076 Tübingen, GERMANY

Camlog Biotechnologies AG
Margarethenstraße 38, 4053 Basel, SWITZERLAND

CAMLOG Management GmbH
Maybachstr. 5, 71299 Wimsheim, GERMANY

**Product
Category(ies):**

High gold dental alloys and solders,
Dental Abutments,
Screws for Dental Abutments,
Healing Caps for Dental Implants, Drills,
Surgery sets and Instruments for Dental Implants
(Class IIa)
Dental Implants, ALTApin Set, Cortico Fix Set
(Class IIb)



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.: 713095093

Valid from: 2017-06-14

Valid until: 2019-06-29

Date, 2017-06-14

Stefan Preiß



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

Page 1 of 1