Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

TÜV SÜD Product Service

EC Certificate

Full Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in class I in sterile conditions, sterilized systems or procedure packs)

No. G1S 076870 0018 Rev. 00

Manufacturer:

ALTATEC GmbH
Maybachstr. 5
71299 Wimsheim
GERMANY

Facility(ies):

CAMLOG Vertrieb GmbH
Maybachstraße 5, 71299 Wimsheim, GERMANY

ALTATEC GmbH
Maybachstr. 5, 71299 Wimsheim, GERMANY

Camlog Biotechnologies GmbH
Margarethenstraße 38, 4053 Basel, SWITZERLAND

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

CAMLOG Management GmbH
Maybachstr. 5, 71299 Wimsheim, GERMANY

Product Category(ies):

Dental Medical Devices supporting insertion, impression taking and bonding

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture in accordance with MDD Annex II. This quality assurance system covers those aspects of manufacture concerned with securing and maintaining sterile conditions of the respective devices / device categories and conforms to the requirements of this Directive. It is subject to periodical surveillance. See also notes overleaf.

Report No.: 713144173

Valid from: 2019-06-30
Valid until: 2024-05-26

Date, 2019-06-28

Stefan Preiß
Head of Certification/Notified Body

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

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

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