

Certificate

Full Quality Assurance System Approval
Annex II excluding (4) of the Directive on Medical
Devices



ECM, Bismarckstr. 106, 52066 Aachen, notified to EC under 0481 hereby declares that an examination of the under mentioned quality assurance system has been carried out following the requirements of annex II excluding (4) of the Directive 93/42/EEC.

This certificate is issued on behalf of:

Manufacturer

TA-Dent GmbH

Moltkestr. 78, 77654 Offenburg, Germany

ECM certifies that the full quality assurance system under which the products listed in annex I to this certificate are manufactured conforms with the requirements of annex II excluding (4) of the Directive 93/42/EEC on medical devices.

This Certificate is only valid for the products mentioned above. Special terms of validity are described in annex I to this certificate.

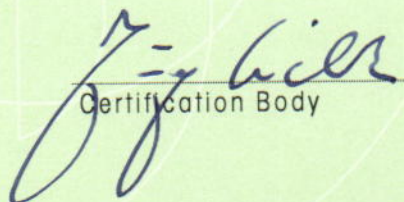
Any substantial changes of the quality assurance system or the listed products which might affect conformity to annex II of the Directive 93/42/EEC have to be notified to ECM and are subject to a separate assessment.

Audit Report Number
678-15-317

Registered under
Z/15/03570E

Valid until
May 17th, 2020

Aachen, May 18th, 2015


Certification Body



Akkreditiert durch
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
ZLG-ZQ-926.94.08



Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
ZLG-BS-240.10.12

This certificate is valid for the hereafter following devices:

Name of product category	Name of individual type	Nomenclature code
dental products	Burs, Dental, Steel	16-669
dental products	Screws	16-055
Non-active implantable products	Prostheses, Dental, Implantable - implants, dental, sterile	16-744
dental products	Prosthetics	/

Special terms of validity:

None.