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

Product Service

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 058693 0015 Rev. 01

Manufacturer:

Being Foshan Medical Equipment Co., Ltd.
No.9 North Park East Road, Shishan Town
Nanhai District
528225 Foshan, Guangdong
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies):

Dental Turbine Handpiece,
Integral Dental Unit,
Dental Electric Motor

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 058693 0015 Rev. 01

Report No.: SH20147EXT01

Valid from: 2021-04-21
Valid until: 2024-05-26

Date, 2021-04-21

Christoph Dicks
Head of Certification/Notified Body

Digitally signed by Ceaicovschi Tudor
Date: 2021.05.20 09:11:21 EEST
Reason: MoldSign Signature
Location: Moldova

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

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