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Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08



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 068173 0020 Rev. 00

Manufacturer:

DIO Corporation

66, Centum seo-ro, Haeundae-gu

Busan 48058

REPUBLIC OF KOREA

**Product Category(ies): Dental Implant Systems and Dental Resins
for 3D Printing of Dental Devices**

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. 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:G10681730020Rev.00

Report No.:

74956756

Valid from:

2021-01-15

Valid until:

2024-05-26

Date,

2021-01-15

Christoph Dicks

Head of Certification/Notified Body