ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
BS-MDR-099

Product Service

EU Quality Management System Certificate (MDR)

Pursuant to Regulation (EU) 2017/745 on Medical Devices, Annex IX Chapters I and III (Class IIa and Class IIb Devices)

No. G10 020326 0062 Rev. 03

Manufacturer:

Institut Straumann AG
Peter Merian-Weg 12
4002 Basel
SWITZERLAND

SRN Manufacturer: CH-MF-000009933

Authorized Representative:

Etkon GmbH
Lochhamer Schlag 6, 82166 Gräfelfing, GERMANY

The Certification Body of TÜV SÜD Product Service GmbH certifies that the manufacturer has established, documented and implemented a quality management system as described in Article 10 (9) of the Regulation (EU) 2017/745 on medical devices. Details on device categories covered by the quality management system are described on the following page(s).
The Report referenced below summarises the result of the assessment and includes reference to relevant CS, harmonized standards and test reports. The conformity assessment has been carried out according to Annex IX Chapter I and III of this regulation with a positive result.
The quality management system assessment was accompanied by the assessment of technical documentation for devices selected on a representative basis.
The certified quality management system is subject to periodical surveillance by TÜV SÜD Product Service GmbH. The surveillance assessment shall also include an assessment of the technical documentation for the device or devices concerned on the basis of further representative samples.
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:G10 020326 0062 Rev. 03

Report No.: 713231132

Preceding Certificate No.: G10 020326 0062 Rev. 02

Valid from: 2022-04-04
Valid until: 2025-06-15

Date of Initial Issuance: 2020-11-26

Christoph Dicks
Head of Certification/Notified Body

Issue date: 2022-04-04

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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No. G10 020326 0062 Rev. 03

Classification:	IIa
Device Group:	L159099 - ODONTOSTOMATOLOGY INSTRUMENTS, REUSABLE - OTHER
Intended Purpose:	-
Classification:	IIa
Device Group:	P010201 - DENTAL IMPLANTS AND ACCESSORIES
Intended Purpose:	-
Classification:	IIa
Device Group:	Q010205 - PRE-FORMED RESINS AND DENTAL CROWNS
Intended Purpose:	-
Classification:	IIa
Device Group:	Q010699 - MATERIALS FOR THE PREPARATION OF CUSTOM-MADE DENTAL DEVICES - OTHER
Intended Purpose:	-
Classification:	IIb
Device Group:	P010201 - DENTAL IMPLANTS AND ACCESSORIES
Intended Purpose:	Straumann® dental implants and abutments are intended for oral implantation to provide a support structure for connected prosthetic devices.
The validity of this certificate depends on conditions and/or is limited to the following:	./.

Revision History:

Rev.	Dated	Report
00	2020-11-26	713176339
01	2021-12-27	713221588
02	2022-03-03	713233299

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