

EU Certificate

Quality Management System REGULATION (EU) 2017/745 on Medical Devices Annex IX Chapters I and III

Registration No.: HZ 1470094-1
Manufacturer: Gebr. Brasseler GmbH & Co. KG
Trophagener Weg 25
32657 Lemgo
Germany

EUDAMED Single
Registration No.: DE-MF-000006446

Products: Products of class IIa:

L090999 ORTHOPAEDIC SURGERY CUTTING
INSTRUMENTS, REUSABLE - OTHER
L159004 ENDODONTIC RASPS AND FILES, REUSABLE
L2499 DERMATOLOGICAL SURGERY INSTRUMENTS,
REUSABLE – OTHER
L26 SURGICAL SCREWDRIVERS, REUSABLE
P091305 BONE SAWS, SINGLE-USE
Q010102 ROOT CANAL FILLING DEVICES
Q010199 CONSERVATIVE DENTISTRY AND
ENDODONTICS DEVICES - OTHER
Q010399 SURGICAL DENTAL DEVICES – OTHER
Q010501 DENTAL BURS AND ABRASIVE DISKS, SINGLE-
USE
Q010502 DENTAL POLISHING CUPS, SINGLE-USE

The Notified Body hereby declares that the requirements of Annex IX, Chapter I of the REGULATION (EU) 2017/745 have been met for the listed products. The above named manufacturer has established and applies a quality management system, which is subject to periodic surveillance, defined by Annex IX, Chapter I, Section 3 of the aforementioned regulation. The requirements of Annex IX, Chapter III are fulfilled.

If class III devices or class IIb implantable devices referred to in the second subparagraph of Article 52(4) are covered by this certificate an EU technical documentation assessment certificate according to Chapter II, Section 4.9 is required before placing them on the market.

Report No.: 1205499-10
Effective date: 2026-04-24
Expiry date: 2031-02-28
Issue date: 2026-04-24



Dipl.-Ing. Ute Frenkert
TÜV Rheinland LGA Products GmbH
Tillystraße 2 · 90431 Nürnberg · Germany

This certificate can be validated on <https://www.certipedia.com>

TÜV Rheinland LGA Products GmbH is a Notified Body according to REGULATION (EU) 2017/745 concerning medical devices with the identification number 0197.

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Products: Products of class IIa:
Q010507 ENDODONTIC INSTRUMENTS (CANAL ENLARGERS, FILES, RASPS, ETC.), SINGLE-USE
V0199 CUTTING DEVICES, SINGLE-USE - OTHER

Products of class I, sterile:

L159004 ENDODONTIC RASPS AND FILES, REUSABLE
Q010199 CONSERVATIVE DENTISTRY AND ENDODONTICS DEVICES - OTHER

The scope of certification is limited to the aspects relating to establishing, securing and maintaining sterile conditions.

Products of class I, reusable surgical instruments:

L159004 ENDODONTIC RASPS AND FILES, REUSABLE
Q010199 CONSERVATIVE DENTISTRY AND ENDODONTICS DEVICES - OTHER

The scope of certification is limited to the aspects relating to the reuse of the device, in particular cleaning, disinfection, sterilization, maintenance and functional testing and the related instructions for use.

Authorized representative(s): N/A



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Certificate history		
Revision:	Description:	Issue date:
6	Re-certification Replaces certificate HZ 1470094-1 Rev. 5 issued 2025-07-17.	2026-02-23
7	Scope extension, Product of class IIa, MDN 1208 (L26)	2026-04-24



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