

EU Certificate

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

Registration No.: HZ 1123799-1

Manufacturer: BEGO Bremer Goldschlägerei
Wilh. Herbst GmbH & Co. KG
Wilhelm-Herbst-Str. 1
28359 Bremen
Germany

EUDAMED Single
Registration No.: DE-MF-000005414

Products:

Products of class IIa:
Q010205 - PRE-FORMED RESINS AND DENTAL CROWNS
Q010601 - DENTAL ALLOYS

Products of class IIb:
Q010601 - DENTAL ALLOYS

Authorized representative(s): N/A

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.: 1152458-260

Effective date: 2026-08-07

Expiry date: 2026-09-10

Issue date: 2026-08-07



Dipl.-Ing. U. 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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Certificate history		
Revision:	Description:	Issue date:
1	First issuance	2022-10-18
2	Scope extension: addition of Q010601	2023-08-23
3	Scope extension: addition of Q010601 under class IIb	2026-08-07

