

EC Certificate
Directive 93/42/EEC Annex II, excluding Section 4
Full Quality Assurance System
Medical Devices

Registration No.: HD 60141590 0001

Report No.: 21248670 017

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

Products: Alloys, SLM Powders, Milling Blanks, Solders, Wires
and Resins for the Dental Field

(see attachment for products included)

Replaces Certificate, Registration No.: HD 60126216 0001


Expiry Date: 2021-08-30

The Notified Body hereby declares that the requirements of Annex II, excluding section 4 of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex II, section 5 of the aforementioned directive. For placing on the market of class III devices covered by this certificate an EC design-examination certificate according to Annex II, section 4 is required.

Effective Date: 2019-08-15

Date: 2018-08-15

Notified Body


Dipl.-Ing. (FH) D. Wiedemuth

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg
TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.

TÜV Rheinland
LGA Products GmbH
Tillystraße 2, 90431 Nürnberg

**Attachment to
Certificate**

Registration No.: HD 60141590 0001
Report No.: 21248670 017

Manufacturer: BEGO Bremer Goldschlägerei
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28359 Bremen
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Products included:

- additives for laser welding, precious metals
- solders, precious metal
- alloys, precious metal
- additives for laser welding, non-precious metals
- solders, non-precious metal
- alloys, non-precious metal
- resins
- SLM Powder
- Milling Blanks

Date: 2019-08-15

Notified Body



Dipl.-Ing. (FH) D. Wiedemuth