

DECLARATION OF CONFORMITY

according to annex II, 93/42/EEC

- Manufacturer: **BEGO Bremer Goldschlägerei
Wilh. Herbst GmbH & Co. KG
Wilhelm-Herbst-Str. 1
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We herewith declare under sole responsibility that the product

- Name of product family: **SLM Powders**
- Name of products: **Mediloy S-Co**
- REF number: **50551**
- Product class: **Class IIb**

meets the relevant requirements of the EC Directive **93/42/EEC** concerning medical devices
and is in conformity with the list of applied standards in the technical documentation.

This declaration of conformity is valid until expiration of the EC-Certificate **93/42/EEC** (Number HD 601126940001 on 30 August 2021.

- Notified body: **TÜV Rheinland LGA Products GmbH
Tillystraße 2
90431 Nürnberg, Germany**

Notified body: **CE 0197**

Bremen, 2017-02-01

Place, Date


Signature
Technical Manager


Signature
Head of Quality Management


Signature
Managing Director

 **BEGO**
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