

DECLARATION OF CONFORMITY

according to annex II without 4, 93/42/EEC

- **Manufacturer:** BEGO Bremer Goldschlägerei
Wilh. Herbst GmbH & Co. KG
Wilhelm-Herbst-Str. 1
28359 Bremen, Germany
T. +49 421 2028-0
F. +49 421 2028-100
www.bego.com

We herewith declare under sole responsibility that the product

- **Name of product family:** **Resins**
- **Name of products:** **Temporary CB**
- **REF number:** FLTCA201, FLTCA301, FLTCB101, FLTCC201
- **Product class:** **Class IIa**

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.

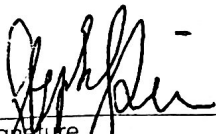
This declaration of conformity is valid until expiration of the EC-Certificate **93/42/EEC** (Number HD 60142369 0001) on 26 May 2024 and only together with the related CoA of a batch.

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

Notified body: **CE 0197**

Bremen, 19.06.2020

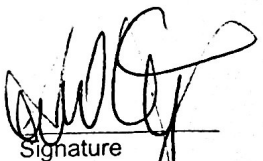
Place, Date



Signature
Chief Development and
Innovation Officer



Signature
Director of Quality Management
and Regulatory Affairs



Signature
Managing Director