

EC Certificate of Conformity

The Notified Body

MEDCERT Zertifizierungs- und Prüfungsgesellschaft für die Medizin GmbH
Pilatuspool 2 – 20355 Hamburg – Germany

herewith certifies that the company:

Bentley InnoMed GmbH
Lotzenäcker 3
72379 Hechingen
Germany

has introduced, applies and maintains a quality assurance system for the products / product categories listed in the appendix.

The compliance of this quality assurance system with the below mentioned requirements of the **Council Directive 93/42/EEC** was verified by an audit:

Annex II without section 4

This certification is subject to surveillance by MEDCERT.

For the placing on the market of class III medical devices covered by this certificate, an additional EC design examination certificate according to Annex II, section 4 of Council Directive 93/42/EEC is required.

Effective date: 2020-06-05

Expiry date: 2024-04-17

Report No.: 7490FS02F

Process No.: QS – 7490

Certificate No.: 7490GB410200605

Hamburg, 2020-06-05



MEDCERT Certification Body
(Dr. Andreas Schich)

The certificate is only valid when provided entirely with all of its pages.
To verify the validity of this certificate, contact info@medcert.de.

MEDCERT Identification Number: 0482



Appendix of EC Certificate of Conformity

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List of products / product categories included in the scope of certificate

- **Coronary stent graft systems**
- **Peripheral vascular stent systems**
- **Peripheral vascular stent graft systems**
- **Aortic stent graft systems**

– End of list –

This appendix is integral part of the above-referenced certificate.
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