

# 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:

**Baat Medical Products B.V.  
F. Hazemeijerstraat 800  
7555 RJ Hengelo, Overijssel  
The Netherlands**

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.

**Effective date: 2020-06-23**


**Expiry date: 2024-01-08**

Report No.: 7512PS02F

Process No.: QS – 7512

Certificate No.: 7512GB410200623

Hamburg, 2020-06-23

  
MEDCERT Certification Body  
(Lorenz Runge)

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

MEDCERT Identification Number: 0482

Form F10010005e EN / Rev. 11 / 2019.11.14



Benannt durch/Designated by  
Zentralstelle der Länder  
für Gesundheitsschutz  
bei Arzneimitteln und  
Medizinprodukten  
[www.zlg.de](http://www.zlg.de)  
ZLG-BS-237.10.15

**Appendix of EC Certificate of Conformity**

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

- **Non-active implantable spinal and intramedullary osteosynthesis devices and related instruments**

– End of list –

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