

# 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:

**OHST Medizintechnik AG**  
**Grünauer Fenn 3**  
**14712 Rathenow**  
**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: 2020-03-19**

**Expiry date: 2024-05-27**

Report No.: 0529FS28F

Process No.: QS – 0529

Certificate No.: 0529GB410200319

Hamburg, 2020-03-19

  
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



## Appendix of EC Certificate of Conformity

Process No.: QS – 0529

Certificate No.: 0529GB410200319

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

- Endoprostheses
- Implants for spine surgery
- Bone screws
- Surgical instruments
- Trial prostheses

– End of list –

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