MEDCERT®

Certification. Medical Only.

# 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:

**ARTOSS GmbH**
**Fischerweg 421**
**18069 Rostock**
**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:** **2020-03-23**
**Expiry date:** **2024-05-27**

Report No.: 3201FS18F
Process No.: QS – 3201
Certificate No.: 3201GB410200323

Hamburg, 2020-03-23

MEDCERT Certification Body
(Markus Bianchi)

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

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-237.10.15

MEDCERT Identification Number: 0482

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## Appendix of EC Certificate of Conformity

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

**Synthetic biodegradeable bone grafting material**

– End of list –

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