MED/CERT®

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:

AQ Solutions GmbH
An der Hasenkaule 10
50354 Hürth
Germany

has introduced 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.

This certificate is valid until 11 November 2023

Report No.: 7510PS02F
Process No.: QS – 7510
Certificate No.: 7510GB410190917

Hamburg, 17 September 2019

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.

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

Process No.: QS – 7510
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List of products / product categories included in the scope of certificate

- **Hip endoprostheses and relating implantation instruments**
- **Knee endoprostheses and relating implantation instruments**
- **Trial prostheses**
- **Tumor and revision system**

– End of list –

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