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:

Heinz Meise GmbH, Medizintechnik
Im Gewerbepark 6
58579 Schalksmühle
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.

Effective date: **2020-10-23**
Expiry date: **2024-05-27**

Report No.: 1203FS26F
Process No.: QS – 1203
Certificate No.: 1203GB410201023A

Hamburg, 2020-10-23

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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Certification. Medical Only.

Appendix of EC Certificate of Conformity

Process No.: QS – 1203
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List of products / product categories included in the scope of certificate

- **Tubing systems**
- **Bloodtubing systems**
- **Plasma containers**
- **Transfer bags**
- **Flushing- and extraction systems**
- **Eye drop application systems**

– End of list –

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