

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:

**Keewell Medical Technology Co., Ltd
Xiananyi Industrial Zone, Pingzhou, Nanhai,
Foshan City, Guangdong 528251
China**

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: 2020-01-27

Expiry date: 2024-05-27

Report No.: 7408IA01F

Process No.: QS – 7408

Certificate No.: 7408GB410200127

Hamburg, 2020-01-27

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.

MEDCERT Identification Number: 0482

Form F10010005e EN / Rev. 11 / 2019.11.14



Benannt durch/Designated by
Zentralstelle der Länder
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bei Arzneimitteln und
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www.zlg.de
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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

- **Blood and infusion warmers**

– End of list –

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