

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

Andreas Fahl Medizintechnik-Vertrieb GmbH
August-Horch-Straße 4a
51149 Köln
Germany

with locations listed in the appendix

has introduced, applies and maintains a quality assurance system
concerning the conformity of the medical devices with the metrological requirements

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 V

This certification is subject to surveillance by MEDCERT.

Effective date: 2020-06-12

Expiry date: 2024-05-27

Report No.: 1320PS26F

Process No.: QS – 1320

Certificate No.: 1320GB416200612

Hamburg, 2020-06-12



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



Appendix of EC Certificate of Conformity

Process No.: QS – 1320

Certificate No.: 1320GB416200612

List of locations included in the scope of certificate

**Andreas Fahl Medizintechnik-Vertrieb GmbH
Am Borsigturm 62
13507 Berlin
Germany**

**Andreas Fahl Medizintechnik-Vertrieb GmbH
Aurbacherstraße 2
81541 München
Germany**

– End of list –

This appendix is integral part of the above-referenced certificate.
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Appendix of EC Certificate of Conformity

Process No.: QS – 1320

Certificate No.: 1320GB416200612

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

Cuff-pressure gauges

– End of list –

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

Andreas Fahl Medizintechnik-Vertrieb GmbH
August-Horch-Straße 4a
51149 Köln
Germany

Hamburg, 2022.03.16

CONFIRMATION

According to the Date of Application (DoA) of the MDR 2017/745 at the 2021.05.27 the Directive 93/42/EEC is no more an active regulation.

For this reason, certificates for Directive 93/42/EEC cannot be issued or changed any more.

As the applied change is not a significant change according to MDCG 2020-3 this change can be accepted, but a revised MDD certificate cannot be issued.

Applied change:

- Deletion of the facility

Andreas Fahl Medizintechnik-Vertrieb GmbH
Aurbacherstraße 2
81541 München
Germany

This confirmation is an attachment to the related certificates

1320DE410200612,
1320GB410200612,
1320DE416200612,
1320GB416200612,
1320DE415200612 and
1320GB415200612.

It confirms that they remain valid including the applied changes and are still applicable.

Best regards



Marcus Harder
Director Certification