EC CERTIFICATE

for the Quality Assurance System

according the directive 93/42/EEC, Annex V

As a notified body of the European Union, DEKRA Certification GmbH certifies, that the company

ERKA. Kallmeyer Medizintechnik GmbH & Co. KG

Im Farchet 15, 83646 Bad Tölz, Germany

applies a quality assurance system for the medical devices listed in the annex according to the directive 93/42/EEC annex V. The approval is based on the result of the re-certification audit report no. 50020-Z5-xx, the decision dated 27.11.2014 is only valid in connection with the successful performance of the annual surveillance audits.

Date of the first certification:	1996-07-29	Date of the last recertification:	2014-11-30
This certificate is valid until:	2017-11-29	Certificate registration No.:	50020-17-06 duplicate

DEKRA Certification GmbH

Stuttgart, 2014-11-27

Notified Body ID-number: 0124

DEKRA Certification GmbH • D-70565 Stuttgart, Handwerkstraße 15

Benannt durch/Designated by
Zentralstelle der Länder für Gesundheitsschutz bei Arzneimitteln und Medizinprodukten
www.zlg.de
ZLG-BS-295.10.02

Lack of fulfilment on conditions as set out in the Certification Agreement may render this certificate invalid

Annex to the Certificate 50020-17-06 dated 27.11.2014

Duplicate

Revision status: 1 Date: 04.12.2014

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Devices/device categories included in the certificate

Class I m:

For the products listed below, the review of the Quality System refers exclusively to the manufacturing steps associated with product conformity and metrological requirements.

- Sphygmomanometers, Aneroide

Class II a:

- Sphygmomanometers, electronic