

CERTIFICATE

for the Quality Assurance System



As a notified body of the European Union (Reg. No. 0124) DEKRA Certification GmbH hereby approves the Quality Assurance System applied for manufacture and final inspection by the company

Hipp Endoskop Service GmbH
Glottertalstraße 10 • 79108 Freiburg, Germany

Approval is based on the decision dated 07.06.2010 and the result of the report no. 50443-Z3-00 and is performed in accordance with the stipulations of

Annex V, Section 3 of the Directive 93/42/EEC

of the Council dated June 14, 1993 governing medical devices. The certification is applicable to the devices specified in the Annex. The devices in question are subjected to testing and examination in accordance with Annex V, Section 3 of the Directive 93/42/EEC. The listed devices may be affixed with the CE marking indicated below.



Date of the first certification: 18.11.1998

This certificate is valid until: 12.11.2014

Date of the last recertification: 07.06.2010

Certificate-registration No.: 50443-17-02
English version (duplicate)

DEKRA Certification GmbH
Stuttgart, 07.06.2010



Akkreditiert durch
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln
und Medizinprodukten
ZLG-ZQ-992.94.16

Annex to the Certificate 50443-17-02 dated 07.06.2010

English version

Revision status: 1

Date: 30.07.2010

Page 1 of 1

Devices/device categories included in the certificate

Class II a:

- rigid endoscopes
- Cannulas, needles and instruments for Endoscopy
- Verres Needles
- Trocar Systems
- Fittings, Adapters, Luer, Stopcocks
- Suction-Irrigation-Instruments
- Suction-Irrigation-Cannulas
- Accessories for Endoscopy
- Elevators



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