



Product Service

# EC-CERTIFICATE

## Full Quality Assurance System

(Annex II, section 3 of the Directive 93/42/EEC on Medical Devices)

No. G1 08 01 40935 010

**Manufacturer:****Allgaier Instrumente GmbH**

Teuchelgrube 6 - 10  
78665 Frittlingen  
GERMANY

**Facility(ies):**

Allgaier Instrumente GmbH  
Teuchelgrube 6 - 10, 78665 Frittlingen, GERMANY

**Product  
Category(ies):**

Orthopaedic Implants, High Frequency Instruments  
and Electrodes, Tracheal cannulae, Special cannulae,  
Sterilization Container Systems, sterilized Scalpels  
and Blades for single use only, Retractors,  
Infrared Ear Thermometer, Insufflators, Suction-/Irrigation  
and Arthroscopy Units, Shaver Systems and Blades,  
Endoscopes and Accessories, endoscopic instruments

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective products / product categories according to Annex II section 3 of the Directive 93/42/EEC on Medical Devices. This quality assurance system conforms to the provisions of this Directive and is subject to periodical surveillance. For marketing of class III products an additional Annex II.4 certificate is mandatory. See also notes overleaf.

**Report No.:**

71332476

**Valid until:**

2009-10-22

**Date,** 2008-02-18

Reiner Krumme



TÜV SÜD Product Service GmbH is Notified Body according to Council Directive 93/42/EEC concerning medical devices with identification no. 0123.

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