

EC Certificate
Directive 93/42/EEC Annex II, excluding Section 4
Full Quality Assurance System
Medical Devices

Registration No.: HD 60107793 0001

Report No.: 21240459 001

Manufacturer: Happersberger otopront GmbH
Langgasse 90
65329 Hohenstein
Deutschland

Products: Medical devices for otorhinolaryngology
(see attachment for products included)

Replaces Approval, Registration No.: HD 60037345 0001

Expiry Date: 2021-01-24

The Notified Body hereby declares that the requirements of Annex II, excluding section 4 of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex II, section 5 of the aforementioned directive. For placing on the market of class III devices covered by this certificate an EC design-examination certificate according to Annex II, section 4 is required.

Effective Date: 2016-01-25

Date: 2016-01-13

Notified Body



Dipl.-Ing. D. Meier

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg

TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.