



Product Service

EC Certificate

Full Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 12 05 31564 011

Manufacturer:**Ulrich AG**

Mövenstraße 12
9015 St. Gallen
SWITZERLAND

**Product
Category(ies):**

Bipolar and monopolar coagulation forceps

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 devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.: 713004952**Valid from:** 2012-08-23**Valid until:** 2017-08-22**Date,** 2012-08-24

Hans-Heiner Junker



TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

Page 1 of 2



Product Service

EC Certificate

Full Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 12 05 31564 011

Facility(ies):

Ulrich AG
Mövenstraße 12, 9015 St. Gallen, SWITZERLAND