



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 13 08 20808 030

**Manufacturer:** MEDICON eG

Gänsäcker 15  
78532 Tuttlingen  
GERMANY

**Facility(ies):**

MEDICON eG  
Gänsäcker 15, 78532 Tuttlingen, GERMANY

**Product Category(ies):**

Active medical devices, non-active medical devices,  
implants and sterile medical devices  
(for detailed information: see attachment)

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.:** 713022688

**Valid from:** 2013-10-13  
**Valid until:** 2018-10-12



**Date,** 2013-10-11

Hans-Heiner Junker

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

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