



Product Service

# EC Certificate

## Production Quality Assurance System

Directive 93/42/EEC on Medical Devices, Annex V  
(Devices in class I with measuring function)

No. G2M 13 07 85278 010

**Manufacturer:** **Carefusion**  
 (Formerly: 1500 Waukegan Road,  
 McGaw Park, IL, 60085)  
 75 North Fairway Drive  
 Vernon Hills IL 60061  
 USA

**EC-Representative:** **CareFusion France 309 S.A.S.**  
 8 bis rue de la Renaissance  
 44110 Châteaubriant  
 FRANCE

**Product Category(ies):** **Class I reusable surgical instruments with measuring function:**  
**Measuring Rods**  
**Measuring Devices for Grafts and Prostheses**  
**Eye Lens Chamber Measuring Devices**  
**Calipers**  
**Rulers**  
**Fenestrometers**  
**Scissors**  
**Markers**  
**Probes**  
**Jigs**  
**Grids**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for the manufacture in accordance with MDD Annex V. This quality assurance system covers those aspects of manufacture concerned with the metrological requirements of the respective devices / device categories and conforms to the requirements of this Directive. It is subject to periodical surveillance. See also notes overleaf.

**Report No.:** NM1306974  
**Valid from:** 2013-09-16  
**Valid until:** 2018-08-03



Hans-Heiner Junker

**Date,** 2013-09-23

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



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**Facility(ies):**

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5 Sunnen Drive, St. Louis, MO 63143, USA

CareFusion  
5175 South Royal Atlanta Drive, Tucker, GA 30084, USA

CareFusion Germany 318 GmbH  
Kantstrasse 33/1 218, 78573 Wurmlingen, GERMANY