



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 17 12 96586 003

**Manufacturer:** CODAN ARGUS AG

Oberneuhofstrasse 10  
6340 Baar  
SWITZERLAND



**Facility(ies):**

CODAN ARGUS AG  
Oberneuhofstrasse 10, 6340 Baar, SWITZERLAND

**Product  
Category(ies):**

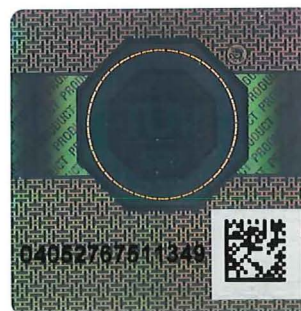
**Infusion Pumps, Syringe Pumps**

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.:** 713122226

**Valid from:** 2017-12-17

**Valid until:** 2022-12-16



**Date,** 2017-12-14

Stefan Preiß

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

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