



Product Service

# EC Certificate

## Production Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex V  
(Devices in class I in sterile conditions, sterilised systems or procedure packs)

No. G2S 15 10 47218 012

**Manufacturer:** **CODAN US Corporation**

3511 West Sunflower Avenue  
Santa Ana CA 92704-6944  
USA



**EC-Representative:** **CODAN Medizinische Geräte GmbH & Co KG**

Stig Husted-Andersen Strasse 11  
23738 Lensahn  
GERMANY

**Product Category(ies):** **Fluid Administration Sets, Enteral Feeding Products**

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

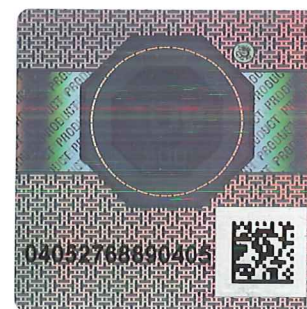
**Report No.:** 72104655

**Valid from:** 2015-12-02

**Valid until:** 2020-12-01

**Date,** 2015-11-23

Hans-Heiner Junker



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

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**Facility(ies):**

CODAN US Corporation

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