ZERTIFIKAT ◆ CERTIFICATE ◆ 認証証書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

Product Service

# EC Certificate

## Production Quality Assurance System

**Directive 93/42/EEC on Medical Devices (MDD), Annex V**
**(Devices in Class IIa, IIb or III)**

**No. G2 16 01 47218 013**

**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

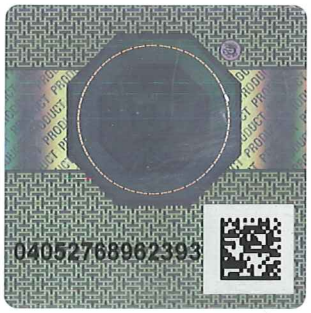
**Product Category(ies):**

**Blood Administration sets**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex V. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class IIb and III devices an additional Annex III certificate is mandatory. See also notes overleaf.

**Report No.:** 72111542

**Valid from:** 2016-05-19
**Valid until:** 2021-05-18




**Date,** 2016-05-19

Stefan Preiß

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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TÜV SÜD Product Service GmbH · Zertifizierstelle · Ridlerstraße 65 · 80339 München · Germany

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