



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 15 09 57451 032

**Manufacturer:****Vogt Medical Vertrieb GmbH**

Rüppurrer Str. 1A Haus B  
76137 Karlsruhe  
GERMANY

**Facility(ies):**

Vogt Medical Vertrieb GmbH  
Rüppurrer Str. 1A Haus B, 76137 Karlsruhe, GERMANY

**Product****Category(ies):**

**Sterile and non sterile I.V. cannulas**  
**Sterile and non sterile disposable epidural needles,**  
**Sterile and non sterile disposable blood bags,**  
**Sterile and non sterile disposable central venous catheter**  
**and kit,**  
**Sterile and non sterile disposable spinal needles,**  
**Sterile and non sterile disposable micro infusion pumps,**  
**Sterile and non sterile disposable silicone catheter Foley**

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.:**

713066362

**Valid from:**

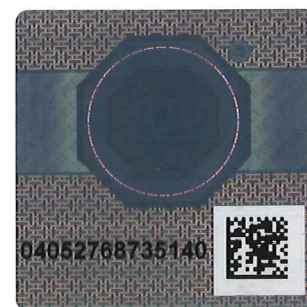
2015-11-23

**Valid until:**

2020-11-22

**Date,** 2015-10-15

Hans-Heiner Junker



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

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