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

TÜV SÜD
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 12 57451 035

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 disposable syringes with needles,
sterile and non sterile disposable scalp vein sets,
sterile and non sterile disposable transfusion sets,
sterile and non sterile disposable infusion sets,
sterile and non sterile disposable surgical gloves,
sterile and non sterile disposable injection needles,
sterile insulin pen needles,
sterile and non sterile disposable laryngeal masks,
sterile and non sterile disposable endotracheal tubes,
sterile and non sterile disposable latex catheters foley,
sterile disposable surgical blades and scalpels

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.: 713095397

Valid from: 2017-02-09
Valid until: 2020-11-09

Date, 2017-02-09

Stefan Preiß

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

TÜV®