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

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 16 10 47326 007

Manufacturer:

IVF HARTMANN AG
Victor-von-Bruns-Strasse 28
8212 Neuhausen am Rheinfall
SWITZERLAND

Facility(ies):

IVF HARTMANN AG
Victor-von-Bruns-Strasse 28, 8212 Neuhausen am Rheinfall,
SWITZERLAND

Product Category(ies):

Compresses, swabs, surgical drapes, primarily wound dressings as well as sterile medical systems for care, wound treatment, therapy, analysis, catheterisation and anaesthesia

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.: 713093072

Valid from: 2017-01-17
Valid until: 2022-01-16

Date, 2016-11-16

Stefan Preiß

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

TÜV®