



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 026269 0006 Rev. 00

Manufacturer

Franz Kalff GmbH

Dechant-Wolfgartenstrasse 85

53881 Euskirchen

GERMANY

Product Category(ies):

**Single-use devices: Bandage materials,
compression bandage, bandages for
general wound treatment, compresses**

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. All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: [www.tuvsud.com/ps-cert?q=cert:G2S 026269 0006 Rev. 00](http://www.tuvsud.com/ps-cert?q=cert:G2S_026269_0006_Rev.00)

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Date, 2021-04-07

Christoph Dicks

Head of Certification/Notified Body