



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 17 09 65765 018**

Manufacturer:**Foosin Medical Supplies Inc., Ltd.**

No.20, Xingshan Road
Weihai Torch Hi-tech Science Park
264210 Weihai, Shandong Province
PEOPLE'S REPUBLIC OF CHINA

**EC-Representative:****MedNet GmbH**

Borkstrasse 10
48163 Muenster
GERMANY

Product**Category(ies):**

non-absorbable surgical suture with or without needle
(PVDF,PTFE,polyester,polypropylene).

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.:

BJ1792607

Valid from:

2017-11-15

Valid until:

2022-08-24

Date, 2017-11-15

Stefan Preis



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

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Facility(ies):

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