



Product Service

CERTIFICATE

No. Q1N 15 06 48773 032

Holder of Certificate: Demophorius Limited



196 Archbishop Makarios III
3030 Limassol
CYPRUS

Facility(ies):

Demophorius Limited
196 Archbishop Makarios III, 3030 Limassol, CYPRUS

Certification Mark:



Scope of Certificate:

Design and Development, Production and Distribution of Sutures, Disposable Medical Devices, Blood collection tubes and blood collection needles, Polypropylene Mesh, Bone Wax, Blood bags, Skin Stapler, Closed Suction Catheter Set

Applied Standard(s):

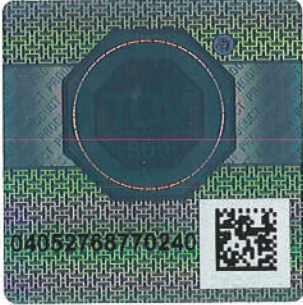
EN ISO 13485:2012 + AC:2012
Medical devices - Quality management systems - Requirements for regulatory purposes (ISO 13485:2003 + Cor. 1:2009)
DIN EN ISO 13485:2012

The Certification Body of TÜV SÜD Product Service GmbH certifies that the company mentioned above has established and is maintaining a quality management system, which meets the requirements of the listed standard(s). See also notes overleaf.

Report No.: 713055218

Valid from: 2015-09-21
Valid until: 2018-07-31

Hans-Heiner Junker



Date, 2015-09-22

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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 08 48773 033**

Manufacturer: **Demophorius Limited**

196 Archbishop Makarios III
3030 Limassol
CYPRUS



Facility(ies):

Demophorius Limited
196 Archbishop Makarios III, 3030 Limassol, CYPRUS

**Product
Category(ies):**

**Non-absorbable and absorbable Sutures,
Sterile Polypropylene Mesh,
Sterile Bone Wax,
Sterile Bloodbags,
Closed Suction Catheter Set**

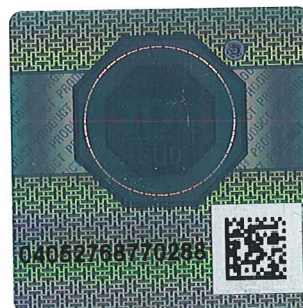
The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms 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.: 713055218

Valid from: 2015-09-21
Valid until: 2017-11-06

Date, 2015-09-22

Hans-Heiner Junker



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

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