



Benannt durch/Designated by  
Zentralstelle der Länder  
für Gesundheitsschutz  
bei Arzneimitteln und  
Medizinprodukten  
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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 074222 0013 Rev. 00**

**Manufacturer:**

**Creagh Medical Ltd**

IDA Business Park

Ballinasloe, Co Galway, H53 K8P4

IRELAND

**Product Category(ies): PTA Catheters for Peripheral Applications  
and Intravascular Microflow Catheters**

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.:** 75944595

**Valid from:** 2020-04-03

**Valid until:** 2022-10-22

**Date,** 2020-04-03

Christoph Dicks

Head of Certification/Notified Body