



Product Service

EC - CERTIFICATE

Full Quality Assurance System

(Annex II, section 3 of the Directive 93/42/EEC on Medical Devices)

No. G1 10 10 39709 683

Manufacturer: **Medtronic Inc.**
710 Medtronic Parkway N.E.
Minneapolis MN 55432
USA

EC-Representative: **Medtronic Ireland**
Parkmore Business Park West
Galway, Ireland
IRELAND

Product Category(ies): **Temporary Occlusion and Aspiration System;
Angioplasty and Angiography Products
(Angiography Catheters, Guiding Catheters,
Diagnostic Catheters, Guidewires, Introducers)**

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 products / product categories according to Annex II section 3 of the Directive 93/42/EEC on Medical Devices. This quality assurance system conforms to the provisions of this Directive and is subject to periodical surveillance. For marketing of class III products an additional Annex II.4 certificate is mandatory. See also notes overleaf.

Report No.: 71374895

Valid until: 2014-12-31

Date, 2010-11-30

Hans-Heiner Junker



TÜV SÜD Product Service GmbH is Notified Body according to Council Directive 93/42/EEC concerning medical devices with identification no. 0123.

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Facility(ies):

Medtronic Vascular
37A Cherry Hill Drive, Danvers MA 01923, USA