



Product Service

EC - CERTIFICATE

Full Quality Assurance System

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

No. G1 09 08 57063 006

Manufacturer: **Invivo Germany, Ein Unternehmen der Philips Medical Systems DMC GmbH**

Hagenower Straße 73
19061 Schwerin
GERMANY

Facility(ies): Invivo Germany, Ein Unternehmen der Philips Medical Systems DMC GmbH
Hagenower Straße 73, 19061 Schwerin, GERMANY

Product Category(ies): **Sterile instruments for interventional radiology and active drilling systems**

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.

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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.

Page 1 of 1