

CERTIFICATE

No. Q1N 16 12 39709 01071

Holder of Certificate: Medtronic Inc.

> 710 Medtronic Parkway N.E. Minneapolis MN 55432

USA

Facility(ies): Medtronic Inc.

7000 Central Avenue N.E., Minneapolis MN

55432, USA

Medtronic Inc.

8200 Coral Sea St., Mounds View MN 55112,

USA

Medtronic Inc., Service Center, Sullivan Lake 800 53rd Avenue NE, Columbia Heights MN

55421, USA

Certification Mark:



Scope of Certificate: Design and development, production and distribution of

Brady and Tachy IPGs, Pacemaker Delivery Systems, Leads and Accessories for Brady and Tachy IPGs, Ablation Systems/ Devices, Diagnostic and Ablation Catheters and their Accessories, Programmers for AIMDs, Application Software (external), External Pacemakers, Software and Hardware Systems for acquisition and management of ECG and cardiac device data,

Implantable Monitoring and Recording Systems, and Service of Programmers for AIMDs, External Pacemakers

and External Patient Monitors

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.:

713091502

Valid from:

2017-01-01

Valid until:

2019-12-31

Date, 2016-12-22

Page 1 of 1

Stefan Preiß





