

EC Design Examination Certificate



No.: I7 02 08 39709 150

according to Annex 2, section 4 of Council Directive 90/385/EEC for active implantable medical devices for

Medtronic Inc.
710 Medtronic Parkway N.E.
Minneapolis MN 55432
USA

with the authorized EC representative:

Medtronic B.V.
Earl Bakkenstraat 10
6422 PJ Heerlen
The Netherlands

Product: Leads for Muscle-/ Neurostimulators and their auxiliary components

Model: see attachment

The product(s) is (are) designed/manufactured in the facility:

Medtronic Gastroenterology and Urology, 4000 Lexington Avenue N., Shoreview, MN 55126, USA

The above mentioned product fulfills the essential requirements of the Directive.

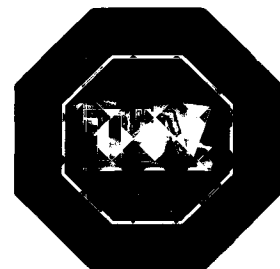
This certificate is based on the examination of the submitted technical design documentation (design dossier). For detailed results of the examination of the design dossier see

Test report no.: 70031812.

This certificate is valid until 2007-08-29.

Released with the above mentioned certificate number by the Certification Body of TÜV PRODUCT SERVICE.

A handwritten signature in black ink, appearing to read 'Pedle'.



Department: MUC AMI / cd / dk
Date: 2002-08-30

TÜV PRODUCT SERVICE GMBH is Notified Body according to Council Directive 90/385/EEC for active implantable medical devices, notified by publication in the Official Journal of the EC No. C 209/8 dated Aug. 3, 1993 with identification no. 0123.

Attachment for Certificate No. I7 02 08 39709 150
from 2002-08-30

**Product: Leads for Muscle-/ Neurostimulators and their
auxiliary components**

Type:	Model:
Unipolar intramuscular stimulation lead	4351
Lead kit for sacral nerve stimulation	3886/3080
Lead kit for sacral nerve stimulation	3092/3966
Lead kit for sacral nerve stimulation	3889/3093
Laryngeal stimulation lead	3090
Quadripolar extension kit for sacral nerve stimulation	3095
Lead introducer	042294
Lead introducer kit	3550-18
Grip-lock anchor accessory kit	3550-17

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