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By Royal Charter

Certificate of Registration

QUALITY MANAGEMENT SYSTEM - ISO 13485:2016 & EN ISO 13485:2016

This is to certify that:

Medtronic Ireland
Parkmore Business Park West
Galway
Ireland

Holds Certificate Number: MD 94974

and operates a Quality Management System which complies with the requirements of ISO 13485:2016 & EN ISO 13485:2016 for the following scope:

The design and manufacture of vascular devices. The manufacture of heart valve delivery and loading systems, heart therapy/pacemaker delivery systems, nonactive implantable/non-implantable medical devices with drug coating/impregnation, catheter systems for renal denervation, venous occlusion systems, atherectomy systems, sterile cardiac ablation catheters and delivery systems, and implantable fixation systems.

For and on behalf of BSI:

Graeme Tunbridge, Senior Vice President Global Regulatory & Quality

Original Registration Date: 2005-03-17
Effective Date: 2024-07-17
Latest Revision Date: 2024-05-29
Expiry Date: 2027-07-16

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Certificate No: MD 94974

Location	Registered Activities
Medtronic Ireland Parkmore Business Park West Galway Ireland	The design and manufacture of vascular devices. The manufacture of heart valve delivery and loading systems, heart therapy/pacemaker delivery systems, nonactive implantable/non-implantable medical devices with drug coating/impregnation, catheter systems for renal denervation, venous occlusion systems, atherectomy systems, sterile cardiac ablation catheters and delivery systems, and implantable fixation systems. The loading & final assembly of transcatheter pacemaker systems. Provision of analytical test services for Medtronic Corporation facilities/sites.
Medtronic, Inc. 710 Medtronic Parkway Minneapolis Minnesota 55432 USA	Administrative Activities.

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Original Registration Date: 2005-03-17
Latest Revision Date: 2024-05-29

Effective Date: 2024-07-17
Expiry Date: 2027-07-16