



CERTIFICATE



Management System as per EN ISO 14001 : 2015

In accordance with TÜV AUSTRIA procedures, it is hereby certified that

MEDTRONIC HELLAS MEDICAL DEVICES S.A **«MEDTRONIC HELLAS S.A.»**

Main Office: 24, Kifisias Avenue, Building B, GR-151 25 Marousi, Attiki, Greece

Diabetes Shop: 2-4, Mesogion Avenue, Athens Tower, GR-115 27 Athens, Greece

Second Office: 9th km Thessaloniki-Nea Moudania, GR-555 35 Thessaloniki, Greece

Applies an Environmental Management System in line with the above Standard for the following Scope

SALES OF MEDICAL DEVICES AND IMPLANTS.

Certificate Registration No.: **20051190002151**

Valid until: 2025-08-07

Initial certification: 2019-08-08

Last certification cycle expiry
date: 2022-08-07

Date of recertification

audit: 2022-07-16

Maria Agapitou
Head of Management Systems & Products Certification Division

Certification Body
at TÜV AUSTRIA

Athens, 2022-08-10

This certification was conducted in accordance with TÜV AUSTRIA auditing and certification procedures and is subject to regular surveillance audits.

TÜV AUSTRIA HELLAS
429, Mesogeion Ave.
GR-153 43 Athens, Greece
www.tuvaustriahellas.gr



Headquarters in Athens bear the responsibility of the Certification decision



CERTIFICATE

The Certification Body of
TÜV SÜD AMERICA INC.

hereby certifies that

Medtronic Energy & Component Center
6800 Shingle Creek Parkway
Brooklyn Center, MN 55430 USA

Has implemented an Environmental Management System in accordance with:

ISO 14001:2015

The scope of this Environmental Management System includes:

**Development, Design and Manufacturing
of Components that Provide, Store, or Transfer
Energy for Implantable Medical Devices**

Certificate Expiry Date: March 2, 2025

Certificate Registration No: 951 13 6301

Issue Date: March 3, 2022

Reissue Date: N/A





Greg Bates
Director Business Assurance America



TÜV SÜD America Inc • 401 Edgewater Place, Suite 500 • Wakefield, MA 01880 USA • www.TUVSUD.com

The validity of this certificate is contingent on the company maintaining its management system to the requirements of the indicated standard and is subject to regular monitoring by TÜV SÜD America.



America

CERTIFICATE

The Certification Body of
TÜV SÜD AMERICA INC.

hereby certifies that

Medtronic PLC Perfusion Systems
7611 Northland Drive
Brooklyn Park, MN 55428 USA

Has implemented an Environmental Management System in accordance with:

ISO 14001:2015

The scope of this Environmental Management System includes:

Design and Manufacturing of Medical Devices

Certificate Expiry Date: February 10, 2025

Certificate Registration No: 951 16 7177

Issue Date: February 11, 2022

Reissue Date: N/A





Greg Bates
Director Business Assurance America



TÜV SÜD America Inc • 401 Edgewater Place, Suite 500 • Wakefield, MA 01880 USA • www.TUVSUD.com

The validity of this certificate is contingent on the company maintaining its management system to the requirements of the indicated standard and is subject to regular monitoring by TÜV SÜD America.



CERTIFICATE

The Certification Body of
TÜV SÜD AMERICA INC.

hereby certifies that

Medtronic
3800 Annapolis Lane
Plymouth, MN 55447 USA

Has implemented an Environmental Management System in accordance with:

ISO 14001:2015

The scope of this Environmental Management System includes:

**Manufacturing of Mechanical Heart Valves
for the Medical Device Industry**

Certificate Expiry Date: June 24, 2024

Certificate Registration No: 951 15 7029

Issue Date: June 25, 2021

Reissue Date: N/A





Greg Bates
Director Business Assurance America





CERTIFICATE

The Certification Body of
TÜV SÜD AMERICA INC.

hereby certifies that

Medtronic, Inc
1851 E. Deere Ave
Santa Ana, CA 92705 USA

Has implemented an Environmental Management System in accordance with:

ISO 14001:2015

The scope of this Environmental Management System includes:

Medical Device Research and Design, Manufacture of Heart Valve Stents, Tissue Processing and Manufacture of Tissue Heart Valves, Manufacture of Thrombectomy Systems and Percutaneous Catheters for the Creation of an Arteriovenous fistula

Certificate Expiry Date: July 18, 2024

Certificate Registration No: 951 15 7089

Issue Date: July 20, 2021

Reissue Date: July 19, 2022





Greg Bates
Director Business Assurance America



TÜV SÜD America Inc • 401 Edgewater Place, Suite 500 • Wakefield, MA 01880 USA • www.TUVSUD.com

The validity of this certificate is contingent on the company maintaining its management system to the requirements of the indicated standard and is subject to regular monitoring by TÜV SÜD America.



NSAI

Certificate of Registration of Environmental Management System to I.S. EN ISO 14001:2015

Medtronic

Parkmore Business Park West
Parkmore
Co. Galway
Ireland

NSAI certifies that the aforementioned company has been assessed and deemed to comply with the provisions of the standard referred to above in respect of:-

The design and manufacture of Cardiology Products.

Approved by:
Stewart Hickey
Head - Business Excellence, NSAI



Registration Number: 14.0381
Original Registration: 16 May 2005
Last amended on: 13 December 2023
Valid from: 13 December 2023
Remains valid to: 15 November 2026

This certificate remains valid on condition that the Approved Environmental Management System is maintained in an adequate and efficacious manner. NSAI is a partner of IQNet – the international certification network (www.iqnet-certification.com)

Partner of:



All valid certifications are listed on NSAI's website – www.nsai.ie. The continued validity of this certificate may be verified under "Certified Company Search"



NSAI (National Standards Authority of Ireland), 1 Swift Square, Northwood, Santry, Dublin 9, Ireland T +353 1 807 3800 E: info@nsai.ie www.nsai.ie

CERTIFICATO



ISO 14001:2015

DEKRA Certification GmbH certifica che l'organizzazione

MEDTRONIC ITALIA S.p.A.

Via Varesina, 162 I - 20156, Milano

per l'area certificata:

Vendita, gestione ordini, magazzino, distribuzione di dispositivi medici, assistenza tecnica, informazione e addestramento a clienti.

ha definito e mantiene attivo un sistema di gestione ambientale secondo la norma sopra citata. La conformità è stata accertata mediante il rapporto di audit n° DK23604867.RA.

N° di registrazione:	170711094/5
Validità certificato precedente:	04.07.2023
Certificato valido da:	05.07.2023
Certificato valido fino al:	04.07.2026



Dr. Rolf Krökel
DEKRA Certification GmbH, Stuttgart, 27.06.2023



Deutsche
Akkreditierungsstelle
D-ZM-16029-01-01

Certificate

for the environmental management system according to

NEN-EN-ISO 14001:2015

including the current version of the underlying SCCM scheme

The certification body TÜV NORD Nederland hereby confirms that the certification took place in accordance with its certification regulations for the organisation

Medtronic B.V.
Earl Bakkenstraat 10
6442 PJ Heerlen
The Netherlands

The management system and the application thereof complies with the requirements of the standard.
The certification is subject to annual evaluation by TÜV NORD Nederland.

Field of application

Receipt, storage, configuration and packaging, order fulfillment and distribution of medical, tissue and pharma products.
Including cleaning of surgical kits and maintenance and calibration within the logistics process.

Registration number 10791-7.1

Start date certificate	18-07-2023
Certificate valid until	04-07-2026
Date of first certificate	04-07-2005
Previous certificate valid until	04-07-2023
Date audit	28-03-2023

Mr. E.W.A.C. Franken
Managing Director



TÜV NORD Nederland B.V.
Ekkersrijt 4401, 5692 DL Son en Breugel
The Netherlands

SCCM
stichting coördinatie
certificatie managementsystemen
voor milieu en gezond en veilig werken

MGMT. SYS.
RVA C 029



Institute of
Quality & Control

CERTIFICATE

NO. 125558

This is to certify that
the Environmental Management System of

Oridion Medical (1987) Ltd.

7 Hamarpe St, Jerusalem, Israel

Was audited by IQC and found to be
in compliance with the requirements of the standard:

ISO 14001 : 2015

This certificate is valid for
the following scope of activities:

**Design & development, manufacture, final inspection and servicing of
Gas sampling lines with and without O2 delivery
Monitors of respiratory monitoring
Madules for respiratory monitoring**

This certificate is valid until: 05.09.2025

Certification cycle will end on: 05.09.2025

Date of first approval: 05.09.2007

This certificate is subject to the continuing satisfactory operation
of the Management System and perloodic auditing by IQC



11.08.2022
Issue date

Nir Halpern, CEO





Certificado del Sistema de Gestión Ambiental



GA-2009/0510

AENOR certifica que la organización

MEDTRONIC IBÉRICA, S.A.

dispone de un sistema de gestión ambiental conforme con la Norma ISO 14001:2015

para las actividades: La distribución y prestación de servicios técnicos, legales y administrativos de productos sanitarios.

que se realiza/n en: OFICINAS: Calle MARIA DE PORTUGAL, 11. 28050 - MADRID
CENTRO DE DISTRIBUCIÓN: PL LA GARENA, PROLOGIS
PARK ALCALA - CL FRANCISCO RABAL, 7 - EDIF. 4. PTA. 1.
28806 - ALCALÁ DE HENARES (MADRID)

Direcciones indicadas en el Anexo

Fecha de primera emisión: 2009-08-17

Fecha de última emisión: 2021-08-17

Fecha de expiración: 2024-08-17



Rafael GARCÍA MEIRO
Director General



THE INTERNATIONAL CERTIFICATION NETWORK

CERTIFICATE

SRAC as an IQNet Partner hereby states that the organization:

MEDTRONIC ROMANIA SRL.

Șoseaua București-Ploiești, nr. 42-44, Clădirea B, aripa B2, et. 2,
Băneasa Business & Technology Park, sector 1, București

for the following scope:

Sale, order management and distribution of active, usual and in vitro diagnostic implantable medical devices, including educating / training clients for the following medical fields: Cardiac ablation, heart rhythm management, cardiovascular surgery, cardiovascular diagnosis, coronary and renal denervation, cranial and spinal interventions, diabetes, ENT, gastrointestinal diagnosis, neuromodulation, neurovascular interventions, patient monitoring, treatment of pelvic and gastric diseases, peripheral and venous, treatment of renal diseases, treatment of respiratory diseases, minimally invasive therapies, medical imaging, robotic surgery. Installation and maintenance of medical devices for the following categories of medical devices: ATI, reusable for surgery, in vitro diagnosis, medical imaging

has implemented and maintains an

Environmental Management System

which fulfils the requirements of the following standard:

ISO 14001 : 2015

Issued on: 2021 - 09 - 15

First issued on: 2021 - 09 - 15

for the validity date, please refer to the original certificate* issued by SRAC

Registration Number: RO - 5778



Alex Stoichitoiu
President of IQNet

eng. Mihaela Cristea
SRAC General Manager



IQNet Partners:**

AENOR Spain AFNOR Certification France APCER Portugal CCC Cyprus CISQ Italy
CQC China CQM China CQS Czech Republic Cro Cert Croatia DQS Holding GmbH Germany FCAV Brazil
FONDONORMA Venezuela ICONTEC Colombia Inspecta Certification Finland INTECO Costa Rica
IRAM Argentina JQA Japan KFQ Korea MIRTEC Greece MSZT Hungary Nemko AS Norway NSAI Ireland PCBC Poland
Quality Austria Austria RR Russia SIGE México SII Israel SIQ Slovenia SIRIM QAS International Malaysia
SQS Switzerland SRAC Romania TEST St Petersburg Russia TSE Turkey Vinçotte Belgium YUQS Serbia
IQNet is represented in the USA by: AFNOR Certification, CISQ, DQS Holding GmbH and NSAI Inc.

* This attestation is directly linked to the IQNet Partner's original certificate and shall not be used as a stand-alone document

** The list of IQNet partners is valid at the time of issue of this certificate. Updated information is available under www.iqnet-certification.com

acreditat pentru
CERTIFICARE



SR EN ISO/CEI 17021-1:2015
CERTIFICAT DE ACREDITARE
SM 004



C E R T I F I C A T

SRAC certifică organizația/ certifies the organisation

MEDTRONIC ROMANIA SRL.

*Șoseaua București-Ploiești, nr. 42-44, Clădirea B, aripa B2, et. 2,
Băneasa Business & Technology Park, sector 1, București*

**pentru următoarele activități/
for the following fields of activities**

Vânzarea, gestionarea comenzilor și distribuția de dispozitive medicale implantabile active, uzuale și pentru diagnostic în vitro, inclusiv educarea / instruirea clienților pentru următoarele domenii medicale: Ablație cardiacă, managementul ritmului cardiac, chirurgie cardiovasculară, diagnosticare cardiovasculară, intervenții coronariene și denervare renală, intervenții craniene și pe coloană vertebrală, diabet, ORL, diagnosticare gastrointestinală, neuromodulație, intervenții neurovasculare, monitorizare pacient, tratamentul afecțiunilor pelvine și gastrice, tratamentul afecțiunilor vasculare periferice și venoase, tratamentul afecțiunilor renale, tratamentul afecțiunilor respiratorii, terapii minim invazive, imagistică medicală, chirurgie robotică. Instalare și mentenanță dispozitive medicale pentru următoarele categorii de dispozitive medicale: A.T.I., reutilizabile pentru chirurgie, diagnostic în vitro, imagistică medicală

Sale, order management and distribution of active, usual and in vitro diagnostic implantable medical devices, including educating / training clients for the following medical fields: Cardiac ablation, heart rhythm management, cardiovascular surgery, cardiovascular diagnosis, coronary and renal denervation, cranial and spinal interventions, diabetes, ENT, gastrointestinal diagnosis, neuromodulation, neurovascular interventions, patient monitoring, treatment of pelvic and gastric diseases, peripheral and venous, treatment of renal diseases, treatment of respiratory diseases, minimally invasive therapies, medical imaging, robotic surgery. Installation and maintenance of medical devices for the following categories of medical devices: ATI, reusable for surgery, in vitro diagnosis, medical imaging

că are implementat și menține un
sistem de management de mediu
conform condițiilor din standardul

which has implemented and maintains an
environmental management system
which fulfils the requirements of the standard

SR EN ISO 14001:2015 (ISO 14001:2015)



Valabilitatea certificatului este condiționată de
efectuarea supravegheților anuale până la data de:



nr. certificat/ certificate registration no. **5778**
data inițială a certificării/ initial certification date **15 septembrie 2021**
data recertificării/ reissuing date -
data ultimei actualizări/ last update -
valabil până la/ valid until **14 septembrie 2024** (cu condiția vizării anuale)
SRAC CERT SRL, Str. Vasile Pârvan Nr. 14, Sector 1, București www.srac.ro

Director General
Ing. Mihaela Cristea

Le système de management de

Medtronic Europe Sàrl Swiss Medtronic Operations (SMO)

Rte de Molliau 31
CH-1131 Tolochenaz



a été audité et certifié selon les exigences de

ISO 14001:2015

Pour les activités suivantes

**Assemblage de dispositifs médicaux actifs implantables
et développement des moyens de production pour cette activité**

Ce certificat est valable du 14 février 2022 au 13 février 2025.
Sa validité est garantie par des audits de surveillance annuels.
L'audit de recertification doit avoir lieu 60 jours ayant la date d'échéance.
Version 5. Certifié depuis février 2013.

L'audit à l'origine de ce certificat a commencé le 1^{er} février 2022.
La validité du certificat précédent était jusqu'au 13 février 2022.

Autorisé par

S. Bieri *D. Willemain*



SGS Société Générale de Surveillance SA
Technoparkstrasse 1 8005 Zurich Switzerland
t +41 (0)44 445-16-80 f +41 (0)44 445-16-88 www.sgs.com



Le système de management de

Medtronic International Trading Sàrl Swiss Headquarters (SHQ)

Rte de Molliau 31
CH-1131 Tolochenaz



a été audité et certifié selon les exigences de

ISO 14001:2015

Pour les activités suivantes

Business support et centre de formation pour l'Europe

Ce certificat est valable du 14 février 2022 au 13 février 2025.
Sa validité est garantie par des audits de surveillance annuels.
L'audit de recertification doit avoir lieu 60 jours avant la date d'échéance.
Version 5. Certifié depuis février 2013.

L'audit à l'origine de ce certificat a commencé le 1^{er} février 2022.
La validité du certificat précédent était jusqu'au 13 février 2022.

Autorisé par

E. Bieri *D. Willemijn*



SGS Société Générale de Surveillance SA
Technoparkstrasse 1 8005 Zurich Switzerland
t +41 (0)44 445-16-80 f +41 (0)44 445-16-88 www.sgs.com

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America

CERTIFICATE

The Certification Body of
TÜV SÜD AMERICA INC.

hereby certifies that

Medtronic Tempe Campus
2343 W. Medtronic Way
Tempe, AZ 85281 USA

Has implemented an Occupational Health and Safety
Management System in accordance with:

ISO 45001:2018

The scope of this Occupational Health and Safety Management System includes:

**Design, Development, and Production
of Microelectronic Components
for Medical Applications**

Certificate Expiry Date: July 29, 2024

Certificate Registration No: 951 21 5381

Issue Date: July 30, 2021

Reissue Date: N/A



Greg Bates
Director Business Assurance America





NSAI

Certificate of Registration of Occupational Health and Safety Management System to I.S. ISO 45001:2018

Medtronic

Parkmore Business Park West
Parkmore
Co. Galway
Ireland

NSAI certifies that the aforementioned company has been assessed and deemed to comply with the provisions of the standard referred to above in respect of:-

The Design and Manufacture of vascular products and the manufacture of heart failure delivery systems

Approved by:
Stewart Hickey
Head - Business Excellence, NSAI



Registration Number: 45.0102
Original Registration: 2 August 2007
Last amended on: 27 July 2022
Valid from: 1 August 2022
Remains valid to: 1 August 2025

This certificate remains valid on condition that the Approved Occupational Health & Safety Management System is maintained in an adequate and efficacious manner.
NSAI is a partner of IQNet - the international certification network (www.iqnet-certification.com)

Partner of:



All valid certifications are listed on NSAI's website - www.nsa.ie. The continued validity of this certificate may be verified under "Certified Company Search"



NSAI (National Standards Authority of Ireland), 1 Swift Square, Northwood, Santry, Dublin 9, Ireland T +353 1 807 3800 www.nsa.ie

Certificate

INTECO has issued an IQNET recognized certificate that the organization:

Medtronic Medical CR S.R.L

Zona Franca Coyoil, edificio B7.6 y B7.5, Alajuela, Costa Rica

has implemented and maintains a

Energy management systems - Requirements with guidance for their use.

for the following scope:

Manufacture of metallic components that are commonly used in spinal surgery, manufactured at the Medtronic Costa Rica plant, located in the Coyoil Free Zone, Coyoil de Alajuela, Costa Rica.

which fulfils the requirements of the following standard INTE/ISO 50001:2018:

INTE/ISO 50001:2018

Issued on: 2024/02/12

First issued on: 2021/03/11

Expires on: 2027/03/11

Registration Number: CR-SGEN-008/202



Alex Stoichitoiu
President of IQNet

Álvaro Torres
Operations director

INTECO 

This attestation is directly linked to the IQNET Member's original certificate and shall not be used as a stand-alone document.

IQNET Members:

AENOR Spain AFNOR Certification France APCER Portugal CCC Cyprus CISO Italy CQC China CQM China CQS Czech Republic
Cro Cert Croatia DQS Holding GmbH Germany EAGLE Certification Group USA FCAV Brazil FONDONORMA Venezuela ICONTEC Colombia ICS
Bosnia and Herzegovina Inspecta Sertifointi Oy Finland INTECO Costa Rica IRAM Argentina JQA Japan KFO Korea LSQA Uruguay MIRTEC Greece
MSZT Hungary Nemko AS Norway NSAI Ireland NYCE-SIGE México PCBC Poland Quality Austria Austria SII Israel SIQ Slovenia SIRIM QAS Interna-
tional Malaysia SQS Switzerland SRAC Romania TSE Turkey YUQS Serbia

*The list of IQNET Members is valid at the time of issue of this certificate. Updated information is available under www.iqnet-certification.com



Certificado de Registro de Empresa Conforme a las exigencias de la norma **INTE/ISO 50001:2018** Sistemas de gestión de la energía. Requisitos con orientación para su uso.

La Asociación Instituto de Normas Técnicas de Costa Rica, INTECO, ubicada en San José, Costa Rica, certifica que la organización:

National Institute of Standardization Association of Costa Rica, INTECO – San José, Costa Rica, certifies that the organization:

Medtronic

Medtronic Medical CR S.R.L

Cumple con los requisitos para el siguiente alcance:

Manufactura de componentes metálicos que se usan comúnmente en cirugías en la columna vertebral, manufacturados en la planta de Medtronic Costa Rica, ubicada en la Zona Franca El Coyol, el Coyol de Alajuela, Costa Rica.

Complies with the requirements for the following scope:

Manufacture of metallic components that are commonly used in spinal surgery, manufactured at the Medtronic Costa Rica plant, located in the Coyol Free Zone, Coyol de Alajuela, Costa Rica.

Que opera en las instalaciones ubicadas en:

which is carried out in
Zona Franca Coyol, El Coyol, Alajuela, Costa Rica.

Director de operaciones

Fecha de emisión: 2021/03/11

Fecha de renovación: 2024/02/12

Fecha de expiración: 2027/03/11

El presente certificado es válido exclusivamente para el proceso descrito, no tiene validez sin su correspondiente alcance de la certificación, donde se indica el servicio, las especificaciones y los documentos normativos aplicables, así como número de registro. En caso de duda siempre prevalecerá el alcance en español. This certificate is valid only for the process described, is not valid without a corresponding scope of certification, indicating the services, specifications and normative documents, and registration number. In case of doubt the spanish written scope will prevail over the english written scope. FS-MC-36. Versión #3. Revisión #A1



Certificate

NSAI has issued an IQNET recognized certificate that the organization:

Medtronic

Parkmore Business Park West, Co. Galway, Ireland

has implemented and maintains an
Energy Management System

for the following scope:

The design and manufacture of vascular products and the manufacture of heart failure delivery systems

which fulfils the requirements of the following standard:

I.S. EN ISO 50001:2018

Issued on: **5 January 2024**
First issued on: **24 November 2008**
Expires on: **25 October 2026**

Registration Number: **IE-93.0005**



Alex Stoichitoiu
President of IQNET



Stewart Hickey
Head - Business Excellence, NSAI



This attestation is directly linked to the IQNET Member's original certificate and shall not be used as a stand-alone document.

IQNET Members*:

AENOR Spain **AFNOR Certification** France **APCER** Portugal **CCC** Cyprus **CISQ** Italy **CQC** China **CQM** China **CQS** Czech Republic
Cro Cert Croatia **DQS Holding GmbH** Germany **EAGLE Certification Group** USA **FCAV** Brazil **FONDONORMA** Venezuela **ICONTEC**
Colombia **ICS** Bosnia and Herzegovina **Inspecta Sertifointi Oy** Finland **INTECO** Costa Rica **IRAM** Argentina **JQA** Japan **KFQ** Korea
LSQA Uruguay **MIRTEC** Greece **MSZT** Hungary **Nemko AS** Norway **NSAI** Ireland **NYCE-SIGE** México **PCBC** Poland **Quality Austria**
Austria **SII** Israel **SIQ** Slovenia **SIRIM QAS International** Malaysia **SQS** Switzerland **SRAC** Romania **TSE** Turkey **YUQS** Serbia

* The list of IQNET Members is valid at the time of issue of this certificate. Updated information is available under www.iqnet-certification.com



NSAI

Certificate of Registration of Energy Management System to I.S. EN ISO 50001:2018

Medtronic

Parkmore Business Park West
Co. Galway
Ireland

NSAI certifies that the aforementioned company has been assessed and deemed to comply with the provisions of the standard referred to above in respect of:-

Energy used in the design and manufacture of vascular products and the manufacture of heart failure delivery systems at this location

Approved by:
Stewart Hickey
Head - Business Excellence, NSAI

Registration Number: 93.0005
Original Registration: 24 November 2008
Last amended on: 8 February 2024
Valid from: 8 February 2024
Remains valid to: 25 October 2026

This certificate remains valid on condition that the Approved Energy Management System is maintained in an adequate and efficacious manner. NSAI is a partner of IQNet the international certification network (www.iqnet-certification.com)

Partner of:



All valid certifications are listed on NSAI's website - www.n sai.ie. The continued validity of this certificate may be verified under "Certified Company Search"



NSAI (National Standards Authority of Ireland), 1 Swift Square, Northwood, Santry, Dublin 9, Ireland T +353 1 807 3800 E: info@nsai.ie www.n sai.ie



BUREAU
VERITAS

MEDTRONIC ITALIA S.P.A.

VIA VARESINA 162 - 20156 MILANO (MI) - Italy

I siti oggetto di certificazione sono in allegato al presente certificato

Bureau Veritas Italia S.p.A. certifica che il sistema di gestione dell'organizzazione sopra indicata è stato valutato e giudicato conforme ai requisiti della norma di sistema di gestione seguente

UNI PdR 125:2022

Sistema di Gestione per la Parità di Genere

Campo di applicazione

Misure per garantire la parità di genere nel contesto lavorativo relativo a Vendita, gestione ordini, distribuzione di dispositivi medici, assistenza tecnica, informazione e addestramento a clienti.

Data della certificazione originale:	05-Febbraio-2024
Data di scadenza precedente ciclo di certificazione:	NA
Data dell'Audit di certificazione / rinnovo:	02-Febbraio-2024
Data d'inizio del presente ciclo di certificazione:	05-Febbraio-2024
Soggetto al continuo e soddisfacente mantenimento del sistema di gestione questo certificato è valido fino al:	04-Febbraio-2027

Certificato Numero: IT329125

Versione: 1

Data di emissione: 05-Febbraio-2024


GLORIA FOCETOLA - Local Technical Manager

ACCREDIA
L'ENTE ITALIANO DI ACCREDITAMENTO

MS N°0009



Indirizzo dell'organismo di certificazione:

Bureau Veritas Italia S.p.A., Viale Monza, 347 - 20126 Milano, Italia

Ulteriori chiarimenti sul campo di applicazione di questo certificato e sui requisiti applicabili della norma del sistema di gestione possono essere ottenuti consultando l'organizzazione.

Per controllare la validità di questo certificato fare doppio click sul QR CODE o scansionarlo con apposita App





BUREAU
VERITAS

Bureau Veritas Certification

Allegato al Certificato di Conformità N° IT329125

MEDTRONIC ITALIA S.P.A.

UNI PdR 125:2022

Sistema di Gestione per la Parità di Genere

Siti oggetto di certificazione

Sito	Indirizzo	Scopo
SEDE OPERATIVA	VIA VARESINA 162 - 20156 MILANO (MI) - Italy	Misure per garantire la parità di genere nel contesto lavorativo relativo a Vendita, gestione ordini, distribuzione di dispositivi medici, assistenza tecnica, informazione e addestramento a clienti.
SITO OPERATIVO	Via Aurelia, 866 - 00165 ROMA (RM) - Italy	
SITO OPERATIVO	Via Campogrande. 43 - 42047 ROLO (RE) - Italy	
SITO OPERATIVO	Via G. Bove 4 - 41037 MIRANDOLA (MO) - Italy	

Certificato Numero: IT329125

Versione: 1

Data di emissione: 05-Febbraio-2024


GLORIA FOCETOLA - Local Technical Manager



MS N°0009

Indirizzo dell'organismo di certificazione:
Bureau Veritas Italia S.p.A., Viale Monza, 347 - 20126 Milano, Italia



Ulteriori chiarimenti sul campo di applicazione di questo certificato e sui requisiti applicabili della norma del sistema di gestione possono essere ottenuti consultando l'organizzazione. Per controllare la validità di questo certificato fare doppio click sul QR CODE o scansionarlo con apposita App

