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ORDIN DE PLATA NR.: 55                                TIP.DOC. 1 :
                                DATA EMITERII:vineri, 7 martie 2:
=====:
PLATITI: 3500-00          LEI: Trei Mii Cinci Sute lei 00 ban :
i                                                                    :
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=====:
PLATITOR:  (R) S.C. "OXIVI    CONTUL DE PLATI/CODUL IBAN      :
T-MED" S.R.L.                MD44ML000000002251729503         :
                                CODUL FISCAL :1007600044280  /   :
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=====:
PRESTATORUL PLATITOR                                CODUL BANCII:
BC"Moldindconbank"S.A. fil."Invest" Chisinau                :MOLDMD2X329:
=====:
BENEFICIAR (R) IMSP SPITAL    CONTUL DE PLATI/CODUL IBAN      :
UL CLINIC MUNICIPAL "SFANTA T MD22ML00000000225166614         :
REIME"                        CODUL FISCAL :1003600152592  /   :
                                                                    :
                                                                    :
=====:
PRESTATORUL BENEFICIAR                                CODUL BANCII:
BC"Moldindconbank"S.A.                                        :MOLDMD2X   :
=====:
DESTINATIA PLATII:Pentru garantia pentru:    TIPUL TRANSFERULUI :
oferta la procedura de achizitie public:      NORMAL/URGENT  :N:
a nr. ocds-b3wdp1-MD-1741075999689 din 1:      :
2.03.2025                                     :
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L.S.                                           :
=====:
                                CODUL TRANZACTIEI:001:          :
                                DATA PRIMIRII:07/03/2025      : SEMNATURILE      :
                                DATA EXECUTARII:              : EMITENTULUI      :
                                :-----:                      :
CONDUCTATOR:Web Kojevnikov Dmitrii
MIIGdAYJKoZIhvcNAQcCoIIGZTCCBmECAQEExCzAJBgUrDgMCGGUAMAsGCSqGSIB3:
DQEHAAACCBH0wggR5MIIDYaADAgECAhNHAADml2rTzDkidh/bAAAAAOaXMA0GCSqG:
SIB3DQEBChUAMCIXIDAeBgNVBAMTF0NFULQxLUNBLU1vbGRpbmRjb25iYW5rMB4X:
DTIzMDMxNjE1MjYyMloXDTI2MDMxNjE1MzYyMlowgbAxCzAJBgNVBAYTAK1EMRAW:
YDVQIQIEwdNb2xkb3ZhMREwDwYDVQQHEWhDaGlzaW5hdTETMBEGA1UEChMKMT3hp
(semnatura electronica)
CONTABIL-SEF:Web Kojevnikov Dmitrii
MIIGdAYJKoZIhvcNAQcCoIIGZTCCBmECAQEExCzAJBgUrDgMCGGUAMAsGCSqGSIB3:
DQEHAAACCBH0wggR5MIIDYaADAgECAhNHAADml2rTzDkidh/bAAAAAOaXMA0GCSqG:
SIB3DQEBChUAMCIXIDAeBgNVBAMTF0NFULQxLUNBLU1vbGRpbmRjb25iYW5rMB4X:
DTIzMDMxNjE1MjYyMloXDTI2MDMxNjE1MzYyMlowgbAxCzAJBgNVBAYTAK1EMRAW:
YDVQIQIEwdNb2xkb3ZhMREwDwYDVQQHEWhDaGlzaW5hdTETMBEGA1UEChMKMT3hp
L.S.                                           (semnatura electronica)
CONDUCTATOR:
(semnatura manuala)
CONTABIL-SEF:
(semnatura manuala)
SEMNATURA PRESTATORUL      L.S.
MOTIVUL REFUZULUI          :-----:
                                : L.S.
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GUVERNUL
REPUBLICII
MOLDOVA



SERVICIUL FISCAL DE STAT



CERTIFICAT

privind lipsa sau existența restanțelor față de bugetul public național

Nr.
№

1431280

Din
От

04.03.2025 14:31



DATE DESPRE CONTRIBUABIL / ИНФОРМАЦИЯ О НАЛОГОПЛАТЕЛЬЩИКЕ

Codul fiscal / Numărul de identificare

Фискальный код / Идентификационный номер

1007600044280

Denumirea

Наименование

Societatea Comercială OXIVIT-MED S.R.L.



ATESTAREA LIPSEI SAU EXISTENȚEI RESTANȚELOR CONFORM DATELOR SISTEMULUI INFORMAȚIONAL AUTOMATIZAT / ПОДТВЕРЖДЕНИЕ ОТСУТСТВИЯ ИЛИ НАЛИЧИЯ ЗАДОЛЖНОСТЕЙ СОГЛАСНО ДАННЫМ ИНФОРМАЦИОННОЙ АВТОМАТИЗИРОВАННОЙ СИСТЕМЫ

La data emiterii prezentului certificat restanța față de bugetul public național constituie

На дату выдачи данной справки задолженность перед национальным публичным бюджетом составляет

0 MDL



VALABIL PÂNĂ LA / ДЕЙСТВИТЕЛЕН ДО

19.03.2025 14:31



Prezentul document este eliberat în temeiul Art. 29, alin. (3) din Legea cu privire la registre nr. 71/2007 și în baza datelor furnizate de Serviciul Fiscal de Stat în Portalul Guvernamental al Cetățeanului și al Unităților de Drept / Справка выдана в соответствие со ст. 29 п. (3) Закона о реестрах № 71/2007 на основании данных, предоставленных Государственной налоговой службой на Портале Правительства Гражданина и Юридических Лиц.

Generat și semnat de Portalul Guvernamental al Cetățeanului și al Unităților de Drept la 04.03.2025 14:31

Prezentul certificat este semnat electronic în conformitate cu Legea nr.124 din 19.05.2022

Сертификат подписан электронной подписью в соответствии с Законом № 124 от 19.05.2022



Certificatul este descărcat din Portalul Guvernamental al Cetățeanului și al Unităților de Drept (mcabinet.gov.md) și este semnat electronic de către posesorul acestui portal și are același valoare juridică ca și documentele eliberate pe suport de hârtie de către organele cu atribuții de administrare fiscală. Verificarea autenticității semnăturii electronice poate fi realizată cu ajutorul Serviciului Guvernamental de Semnătură Electronică (msign.gov.md)

Сертификат скачен с Правительственного Портала Гражданина и Юридических Лиц (mcabinet.gov.md) и подписан электронной подписью владельца портала и имеет такую же юридическую силу, как и документы выдаваемые на бумаге органами налоговой администрации. Проверку подлинности электронной подписи можно осуществить с помощью Государственной Службой Электронной Подписью (msign.gov.md)

REPUBLICA



MOLDOVA

CERTIFICAT DE ÎNREGISTRARE

Societatea Comercială "OXIVIT-MED" S.R.L.
ESTE ÎNREGISTRATĂ LA CAMERA ÎNREGISTRĂRII DE STAT

Numărul de identificare de stat - codul fiscal
1007600044280

Data înregistrării

30.07.2007

Data eliberării

30.07.2007

Bordeianu Tatiana, registrator de stat

*Funcția, numele, prenumele persoanei
care a eliberat certificatul*

semnătura

MD 0067985





AGENȚIA SERVICII PUBLICE

Departamentul înregistrare și licențiere a unităților de drept

EXTRAS din Registrul de stat al persoanelor juridice

Nr. 531861 data 19.09.2023

Denumirea completă: **Societatea Comercială "OXIVIT-MED" S.R.L.**

Denumirea prescurtată: **S.C. "OXIVIT-MED" S.R.L.**

Forma juridică de organizare: **Societate cu răspundere limitată,**

Numărul de identificare de stat și codul fiscal (IDNO): **1007600044280**

Data înregistrării de stat: **30.07.2007**

Sediul: **MD-2032, bd . Decebal, 82, ap.(of.) 90, mun. Chișinău, Republica Moldova.**

Obiectul principal de activitate:

- 1. Fabricarea, comercializarea, asistența tehnică, repararea și verificarea articolelor de tehnică și optică medicală**
- 2. Comerțul cu ridicata al parfumurilor și produselor cosmetice**
- 3. Comerțul cu amănuntul al produselor cosmetice și de parfumerie, articolelor de toaletă**
- 4. Intermedieri pentru vânzarea unui asortiment larg de mărfuri**
- 5. Alte tipuri de comerț cu amănuntul în magazine nespecializate**
- 6. Alte tipuri de comerț cu ridicata**
- 7. Închirierea altor mașini și echipamente**

Capitalul social: **5400 lei,**

Administrator: **KOJEVNIKOV DMITRII, IDNP 0972305012362,**

Asociații:

1. **KOJEVNIKOV DMITRII, IDNP 0972305012362, cota 5400 lei, ce constituie 100%**

Beneficiar efectiv:

1.1. **KOJEVNIKOV DMITRII, IDNP 0972305012362**

Prezentul extras este eliberat în temeiul art.34 al Legii nr.220-XVI din 19 octombrie 2007 privind înregistrarea de stat a persoanelor juridice și a întreprinzătorilor individuali și confirmă datele din Registrul de stat la data de: **19.09.2023.**

**Registrator în domeniul
înregistrării de stat**

Digitally signed by Rusu Diana
Date: 2023.09.19 11:22:47 EEST
Reason: MoldSign Signature
Location: Moldova



Rusu Diana



EB 0461498

OXIVIT MED

c/f: 1007600044280; adresa: str. Decebal 82-90, or. Chișinău, Republica Moldova
telefon: + 373 22 808002; fax: + 373 22 808003
web: www.oxivit-med.com; e-mail: info@oxivit-med.com

Lista fondatorilor companiei SRL „Oxivit-Med”

Nr.	Numele, Prenumele	Codul Personal
1	Kojevnikov Dmitrii	0972305012362

ClosureFast™ Endovenous Radiofrequency Ablation Catheter EU MDR Declaration of Conformity

D00372256

Revision C

Page 1 of 5

Form

Medtronic

EU MDR Declaration of Conformity (DoC)

Manufacturer:	Medtronic, Inc. 710 Medtronic Parkway Minneapolis MN 55432 USA
Manufacturer SRN:	US-MF-000019977
Authorized Representative:	Medtronic B.V. Earl Bakkenstraat 10 6422 PJ Heerlen The Netherlands
Authorized Representative SRN:	NL-AR-000006050
Notified Body:	BSI Group The Netherlands B.V. Say Building, John M. Keynesplein 9 1066 EP Amsterdam The Netherlands Notified Body Number: 2797
Conformity Assessment Certificate(s):	MDR 734659 EU Quality Management System Certificate
Conformity Assessment Route:	Annex IX, Conformity Assessment Based on a Quality Management System
Risk Class:	Class IIb
Classification rule:	Annex VIII Rule 9
Intended purpose:	The ClosureFast catheter is intended for endovascular coagulation of blood vessels in patients with superficial vein reflux.

ClosureFast™ Endovenous Radiofrequency Ablation Catheter EU MDR Declaration of Conformity

D00372256

Revision C

Page 2 of 5

Form

Medtronic

Statement:

We, Medtronic, Inc. hereby declare under our sole responsibility that the product(s) specified herein conform to EU Medical Device Regulation 2017/745 and relevant Union Legislation that provides for the issuing of an EU Declaration of Conformity.

Medtronic, Inc. hereby declares that the above-mentioned products are compliant with the Restriction of Certain Hazardous Substances in Electrical and Electronic Equipment, described in "Directive 2011/65/EU of the European parliament and of the Council of 8 June 2011", commonly known as the EU RoHS Directive

Union Legislation	Declaration of Conformity Document Number
RoHS directive (EU) 2011/65 (RoHS II)	D00372256 (included with this DOC)

ClosureFast™ Endovenous Radiofrequency Ablation Catheter EU MDR Declaration of Conformity

D00372256

Revision C

Page 3 of 5

Form

Medtronic

Place: Plymouth, MN

Name: Laura Moen-Ftacek

Title: Regulatory Affairs Director

Signature: Laura Moen-Ftacek

Date: 20 July 2023

ClosureFast™ Endovenous Radiofrequency Ablation Catheter EU MDR Declaration of Conformity

D00372256

Revision C

Page 4 of 5

Form

Medtronic

Products Covered

Product Name	Medtronic Product Identifier	Basic UDI-DI	Optional: Additional nomenclature identifier (e.g. EMDN, GMDN)
	CFN		
ClosureFast™ EndoVenous Radio frequency Ablation (RFA) Catheter	CF7-7-60	0763000B00004057C	C01901902
	CF7-7-100		
	CF7-3-60		
	CF6-8-60		
	CF6-8-100		

ClosureFast™ Endovenous Radiofrequency Ablation Catheter EU MDR Declaration of Conformity

D00372256

Revision C

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Form

Medtronic

Common Specification(s)

The following common specifications were used to demonstrate conformity:

Number	Date of Issue	Title
Not Applicable		

EU Quality Management System Certificate

Regulation (EU) 2017/745, Annex IX Chapter I and III

MDR 734659 R000

Manufacturer: Medtronic, Inc.

Address:

710 Medtronic Parkway
Minneapolis
MN
55432
USA

Single Registration Number: US-MF-000019977

EU Authorised Representative: Medtronic B.V.

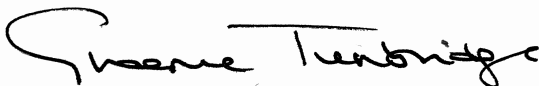
Address:

Earl Bakkenstraat 10
6422 PJ Heerlen
The Netherlands

Scope: See attached **Device Schedule**

On the basis of our examination of the quality system in accordance with Regulation (EU) 2017/745, Annex IX Chapter I and III, the quality system meets the requirements of the Regulation. For the placing on the market of Class III devices, and Class IIb implantable devices that are not considered well-established technologies as specified in Article 52(4) an additional Annex IX Chapter II certificate is required.

For and on behalf of BSI, a Notified Body for the above Regulation (Notified Body Number 2797):



Graeme Tunbridge, Senior Vice President Medical Devices

First Issue Date: **2021-06-09**

Current Issue Date: **2024-03-07**

Starting Validity Date: **2024-03-07**

Expiry Date: **2026-06-08**

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EU Quality Management System Certificate

Regulation (EU) 2017/745, Annex IX Chapter I and III

MDR 734659 R000

Device Schedule: Class III and Class IIb devices

Class III, Implantable	Intended purpose
VenaSeal™ Closure System	See MDR 734883
Class IIb, Implantable	Intended purpose
Abre Venous Self-expanding Stent System	See MDR 734811
Class IIb	Intended purpose
Radiofrequency Venous Ablation Instruments	The Radiofrequency Catheter is intended to be used in conjunction with the Radiofrequency Generator in order to occlude the treated vessel via a fibrotic seal. The Radiofrequency Stylet is intended for use in vessel and tissue coagulation including treatment of incompetent (i.e., refluxing) perforator and tributary veins.
Radiofrequency Generators	The Radiofrequency Generator is intended to provide treatment information and to provide RF energy to the Radiofrequency Catheter or Radiofrequency Stylet when they are used for vessel and tissue coagulation. The Radiofrequency Generator is used with radiofrequency catheters intended for vessel and tissue coagulation.

First Issue Date: **2021-06-09**

Current Issue Date: **2024-03-07**

Starting Validity Date: **2024-03-07**

Expiry Date: **2026-06-08**

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EU Quality Management System Certificate

Regulation (EU) 2017/745, Annex IX Chapter I and III

MDR 734659 R000

Certificate History

(References to applicable Common Specifications, Harmonized Standards complied with, and the relevant test and audit reports that support any of the below certificate changes may be requested from Certificate.Verification@bsigroup.com)

Date	Reference number	Action
2021-06-09	3277806	Issued
2022-02-21	3615666	Supplemented – Addition of ClosureFast RFA Catheter and VenaSeal Closure System to device schedule. Amended – Addition of subcontractors.
2022-05-13	3658991	Supplemented – Addition of ClosureRFG Radiofrequency Generator. Amended – Addition of subcontractors. Addition of manufacturer's SRN. Editorial correction to legal manufacturing address from Minnesota to MN.
2023-02-01	3748363	Amended – administrative update to the device grouping of Class IIb devices in the product table. Removed Critical Subcontractor and Crucial Supplier information. Removed subcontractor information from Certificate History.
2023-03-17	3872264	Amended – Change of VenaSeal™ Closure System classification from Class IIb, Implantable to Class III, Implantable
2023-12-01	30039074	Amended – addition of alternate sterilization subcontractor and modified EO cycle parameters for Radiofrequency Venous Ablation Instruments.
Current	30107348	Amended – Change to subcontractor address

First Issue Date: **2021-06-09**

Current Issue Date: **2024-03-07**

Starting Validity Date: **2024-03-07**

Expiry Date: **2026-06-08**

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Page 3 of 3

Validity of this certificate is conditional on the Manufacturer's quality system being maintained to the requirements of the Regulation as demonstrated through the required surveillance activities of the Notified Body.

This certificate was issued electronically and is bound by the conditions of the contract.

NB Contact: BSI Group The Netherlands B.V., Say Building, John M. Keynesplein 9, 1066 EP, Amsterdam, Netherlands. Tel: + 31 (0) 20 346 07 80
Corporate Contact: BSI Group Assurance Limited, registered in England under number 05435540 at 389 Chiswick High Road, London, W4 4AL, UK.
A Member of the BSI Group of Companies.