ORDIN DE PLATA NR.: 11 TIP.DOC. 1
DATA EMITERII:vineri, 24 ianuari

PLATITI: 2500-00 LEI: Doua Mii Cinci Sute lei 00 bani

PLATITOR: (R) S.C. "OXIVIT-MED" S.R.L.
CONTUL DE PLATI/CODUL IBAN
MD44ML000000002251729503
CODUL FISCAL :1007600044280 /

PRESTATORUL PLATITOR
BC"Moldindconbank"S.A. fil."Invest" Chisinau
CODUL BANCII:
:MOLDMD2X329:

BENEFICIAR (R) I.M.S.P. "Spitalul Clinic Republican Timofei Mosneaga"
CONTUL DE PLATI/CODUL IBAN
MD57MO2251ASV96476607100
CODUL FISCAL :1003600150783 /

PRESTATORUL BENEFICIAR
Mobiasbanca-OTP Group S.A.
CODUL BANCII:
:MOBBMD22

DESTINATIA PLATII:Pentru garantia pentru oferta la procedura de achizi?ie publica nr. ocds-b3wdp1-MD-1735287369911 din 27.01.2025

TIPUL TRANSFERULUI
NORMAL/URGENT :N:

L.S.

CODUL TRANZACTIEI:001:
DATA PRIMIRII:24/01/2025
DATA EXECUTARII:

SEMNATURILE EMITENTULUI

CONDUCATOR:Web Kojevnikov Dmitrii

MIIGdAYJKoZIhvcNAQcCoIIGZTCCBmECAQExCzAJBgUrDgMCGgUAMAsGCSqGSIb3:
DQEHAaCCBH0wggR5MIIDYaADAgECAhNHAADml2rTzDkidh/bAAAAAOaXMA0GCSqG:
SIb3DQEBCwUAMCIxIDAeBgNVBAMTF0NFUlQxLUNBLU1vbGRpbmRjb25iYW5rMB4X:
DTIzMDMxNjE1MjYyMloXDTI2MDMxNjE1MzYyMlowgbAxCzAJBgNVBAYTAk1EMRAw:
YDVQQIEwdNb2xkb3ZhMREwDwYDVQQHEwhDaGlzaW5hdTETMBEGA1UEChMKT3hp

(semnatura electronica)

CONTABIL-SEF:Web Kojevnikov Dmitrii

MIIGdAYJKoZIhvcNAQcCoIIGZTCCBmECAQExCzAJBgUrDgMCGgUAMAsGCSqGSIb3:
DQEHAaCCBH0wggR5MIIDYaADAgECAhNHAADml2rTzDkidh/bAAAAAOaXMA0GCSqG:
SIb3DQEBCwUAMCIxIDAeBgNVBAMTF0NFUlQxLUNBLU1vbGRpbmRjb25iYW5rMB4X:
DTIzMDMxNjE1MjYyMloXDTI2MDMxNjE1MzYyMlowgbAxCzAJBgNVBAYTAk1EMRAw:
YDVQQIEwdNb2xkb3ZhMREwDwYDVQQHEwhDaGlzaW5hdTETMBEGA1UEChMKT3hp

L.S. (semnatura electronica)

CONDUCATOR: _____
(semnatura manuala)

CONTABIL-SEF: _____
(semnatura manuala)

SEMNATURA PRESTATORUL L.S.

Digitally signed by Kojevnikov Dmitrii
Date: 2025.01.24 14:13:44 EET
Reason: MoldSign Signature
Location: Moldova

MOTIVUL REFUZULUI

MOLDOVA EUROPEANĂ

MOBIAS BANCA

Nr. 12101-304

18.03.2016

CERTIFICAT
PRIVIND EXISTENTA CONTURILOR CURENTE

Prin prezentul, **BC „Mobiasbancă – Groupe Societe Generale" S.A.**, codul băncii (BIC): **MOBBMD22**, confirmă că compania **OXIVIT-MED SRL**, cod fiscal (IDNO) **1007600044280,** deține următoarele conturi curente la BC "Mobiasbancă-Groupe Societe Generale" S.A., Filiala. 1 Stejaur :

1. **MDL - 2224710SV23488147100; IBAN- MD09MO2224ASV23488147100**
2. **EUR - 2224710SV22227957100; IBAN- MD17MO2224ASV22227957100**
3. **USD - 2224710SV22214937100; IBAN- MD86MO2224ASV22214937100**

Certificatul este emis în baza cererii întreprinderii: Oxivit-Med SRL.

L.Ș.

Dumitru Popa
Director filială „Stejaur"

Executor : Mariana Guzun
Tel: 022 812 614

Filiala Nr. 1 „Stejaur"
Bd. Ştefan cel Mare şi Sfânt 196
MD-2004, Chişinău, Moldova
Cod MOBBMD22
Cont de corespondenţă 35213892
la Centrul de Decontări al BNM

Tel. +373 22 81 26 15
Fax. +373 22 81 26 15
www.mobiasbanca.md

BC „Mobiasbancă – Groupe Société Générale" SA
Capital Social: 100 000 000 MDL
Număr de înregistrare de stat - 1002600006089
Sediul Central:
bd. Ştefan cel Mare şi Sfânt 81a
MD-2012, Chişinău, Moldova

GROUPE SOCIETE GENERALE

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

CAMERA ÎNREGISTRĂRII DE STAT A REPUBLICII MOLDOVA L.Ș.

MD 0067985

AGENŢIA SERVICII PUBLICE

Departamentul înregistrare şi licenţiere a unităţilor 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 şi codul fiscal (IDNO): **1007600044280**
Data înregistrării de stat: **30.07.2007**
Sediul: **MD-2032, bd . Decebal, 82, ap.(of.) 90, mun. Chişinău, Republica Moldova.**
Obiectul principal de activitate:
1. Fabricarea, comercializarea, asistenţa tehnică, repararea şi verificarea articolelor de tehnică şi optică medicală
2. Comerţul cu ridicata al parfumurilor şi produselor cosmetice
3. Comerţul cu amănuntul al produselor cosmetice şi de parfumerie, articolelor de toaletă
4. Intermedieri pentru vînzarea unui asortiment larg de mărfuri
5. Alte tipuri de comerţ cu amănuntul în magazine nespecializate
6. Alte tipuri de comerţ cu ridicata
7. Închirierea altor maşini şi echipamente

Capitalul social: **5400** lei,

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

Asociaţii:
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 şi a întreprinzătorilor individuali şi confirmă datele din Registrul de stat la data de: **19.09.2023**.

Registrator în domeniul înregistrării de stat **Rusu Diana**

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

EB 0461498

Date cu caracter personal

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

GMED
GROUPE LNE

ATTESTATION / *CERTIFICATE* **N° 8858 rev. 22**
Délivrée à Paris le 06 mai 2021
Issued in Paris on May 6th, 2021

ATTESTATION CE / *EC CERTIFICATE*

Approbation du Système Complet d'assurance Qualité/ *Approval of full Quality Assurance System*
ANNEXE II excluant le point 4 Directive 93/42/CEE relative aux dispositifs médicaux
ANNEX II excluding section 4 Directive 93/42/EEC concerning medical devices
Pour les dispositifs de classe III, un certificat CE de conception est requis
For class III devices, a EC design certificate is required

Fabricant / *Manufacturer*

MEDTRONIC MINIMED
18000 Devonshire street
NORTHRIDGE, CA 91325 UNITED STATES

Catégorie du(des) dispositif(s) / *Device(s) category*

Système de mesure en continu du glucose

Continuous glucose monitoring system

Voir document complémentaire GMED / See GMED additional document
n° 37095

GMED atteste qu'à l'examen des résultats figurant dans le rapport référencé P600181 - P600985, le système d'assurance qualité - pour la conception, la production et le contrôle final - des dispositifs médicaux énumérés ci-dessus est conforme aux exigences de l'annexe II excluant le point 4 de la Directive 93/42/CEE.

GMED certifies that, on the basis of the results contained in the file referenced P600181 - P600985, the quality system - for design, manufacturing, and final inspection - of medical devices listed here above complies with the requirements of the Directive 93/42/EEC, annex II excluding section 4.

La validité du présent certificat est soumise à une vérification périodique ou imprévue
The validity of the certificate is subject to periodic or unexpected verification

Début de validité / *Effective date :* **May 6th, 2021 (included)**
Valable jusqu'au / *Expiry date :* **May 26th, 2024 (included)**

DocuSigned by:
Lionel DREUX
A1D80E08C60D47A...

GMED SAS GMED SAS GMED GROUPE LNE

Lionel DREUX
Certification Director

GMED_b2p3_new2020-V0-04-

GMED - 8858 rev. 22
Modifie le certificat 8858-21

GMED
GROUPE LNE

Document complémentaire GMED n° 37095 rev. 1
GMED additional document n° 37095 rev. 1
Dossier(s) / *File(s)* N° P600181 - P600985
Délivré à Paris le 06/05/2021
Issued in Paris on 05/06/2021

Ce document complémentaire GMED n° 37095 rev. 1 atteste de la validité du certificat CE n° 8858 rev. 22 au regard des informations listées ci-dessous.

This GMED additional document n° 37095 rev. 1 attests to the validity of CE certificate n° 8858 rev. 22 with regard to the information listed below.

Fabricant / *Manufacturer:* **MEDTRONIC MINIMED**
18000 Devonshire street
NORTHRIDGE, CA 91325, UNITED STATES

Identification des dispositifs / *Identification of devices*

Désignation du dispositif / Accessoires marqués CE *Device designation / CE marked accessories*	**Réf commerciale du dispositif ou code article** *Device commercial reference or article code*	**Classe du DM** *MD class*
CGM Transmitters	**MiniLink REAL-Time Transmitter** (MMT-7703, MMT-7735, MMT-7737)	**IIa**
	MiniLink REAL-Time Transmitter System (MMT-7774, MMT-7725)	
	Guardian 2 Link Transmitter (MMT-7731, MMT-7738)	
	Guardian 2 Link Transmitter System (MMT-7730, MMT-7775)	
	Guardian Connect Transmitter (MMT-7821, MMT-7821L)	
	Guardian Connect Transmitter kit (MMT-7820, MMT-7820L)	
	Guardian Link (3) Transmitter (MMT-7811, MMT-7911)	
	Guardian Link (3) Transmitter Kit (MMT-7810, MMT-7910)	
	iPro2 Continuous Glucose Monitoring Digital Recorder (MMT-7741)	
	iPro2 Continuous Glucose Monitoring Digital Recorder System (MMT-7745)	
	Guardian 4 Transmitter (MMT-7841, MMT-7841Q)	
Glucose Sensors	**Enlite Sensor** (MMT-7008A, MMT-7008B)	**IIb**
	Guardian Sensor (3) (MMT-7020C, MMT-7020D)	
	Guardian 4 Sensor (MMT-7040C, MMT-7040D, MMT-7040QC, MMT-7040QD)	

DocuSigned by:
Lionel DREUX
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GMED 0459

GMED - 37095 rev.1

Lionel DREUX
Certification Director

Désignation du dispositif / Accessoires marqués CE *Device designation / CE marked accessories*	**Réf commerciale du dispositif ou code article** *Device commercial reference or article code*	**Classe du DM** *MD class*
CGM Applications	**Guardian Connect App** (CSS7200) (v3), (CSS7201) (v3)	**IIa**
	CareLink Software for Professional CGM Systems (MMT-7340) (v2)	
	Guardian Application (MMT-8200, MMT-8201)	

Sites couverts et Activités / *Locations and Activities*

Site / *Location*	**Activités** / *Activities*
MEDTRONIC MINIMED **18000 Devonshire street** **NORTHRIDGE, CA 91325** **UNITED STATES**	**Conception, fabrication et contrôle final** *Design, manufacture and final control*

DocuSigned by:
Lionel DREUX
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GMED 0459

GMED - 37095 rev.1

Lionel DREUX
Certification Director

EC DECLARATION OF CONFORMITY

DoC No:	2021-002-01
Manufacturer:	Medtronic MiniMed 18000 Devonshire Street Northridge CA, 91325 USA
EC Representative:	Medtronic BV Earl Bakkenstraat 10 6422 PJ Heerlen The Netherlands
Conformity Assessment Procedure: Annex II, except section 4	MDD 93/42/EEC Medical Devices Directive
EC Quality System Certificate: (Annex II, excluding section 4)	Certificate No. 8858 (Expiry: May 26th, 2024)
Notified Body:	GMED 0459 1 rue Gaston Boissier 75015 PARIS FRANCE
Description of Device Concerned:	**Guardian™ 4 Sensor** MMT-7040C (5 pack) MMT-7040D (1 pack) MMT-7040QC (5 pack) MMT-7040QD (1 pack)
Classification:	Class IIb (Rule 8, Annex IX)
GMDN Code:	59016
Technical File Reference:	10912317DOC_E
Conformity with the following standard(s) or other normative document(s)	Refer to Essential Requirement Matrix 9181135DOC_X

Statement:
I, the undersigned, hereby declare on behalf of Medtronic MiniMed and under its sole responsibility, that the Medical Device(s) specified above and provided with CE marking, conform with the Essential Requirements of the EC Directive 93/42/EEC Medical Device Directive. All supporting documentation is retained under the premises of the manufacturer.

G George

Geena George
Sr. Regulatory Affairs Manager
Medtronic MiniMed
18000 Devonshire Street
Northridge CA, 91325, USA

March 12, 2021
Date

Valid from signature date

Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

Product Service

EC Certificate

Full Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 039709 1263 Rev. 00

Manufacturer:

Medtronic, Inc.
710 Medtronic Parkway
Minneapolis MN 55432
USA

Product Category(ies):

- **Autotransfusion Systems and Associated Disposables**
- **Centrifugal Blood Pumps**
- **Bio-Console Drive Units**
- **Flow Monitoring Systems**
- **Bio-Cal Blood Temperature Controller**
- **Temperature Monitoring Systems and Associated Disposables**
- **Blood Monitoring Systems**
- **Cardioplegia Delivery Systems**
- **Disposable Blood Handling Devices used for Open Heart Surgery**
- **Arterial Filters**
- **Oxygenators including Heat Exchangers, with and without Cardiotomy Reservoirs**
- **Cardiotomy Venous Reservoirs**
- **Venous Reservoir Bags**
- **Perfusion Equipment and Disposable Perfusion Devices**
- **Disposable Medical Devices for Drainage Systems**
- **Disposable Medical Devices for use in Cardiopulmonary Surgery: Cardioplegia, Cannulae, Venting, Suction**
- **Pressure Display System & related accessories of class IIa**
- **Tissue Positioning/Stabilizing Devices**
- **Surgical Site Clearing Devices**
- **Intravascular Shunts**
- **Surgical Retractors**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

Benannt durch/Designated by
Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
www.zlg.de
ZLG-BS-244.10.08

Product Service

EC Certificate

Full Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 039709 1263 Rev. 00

inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.: 72150396

Valid from: 2020-02-12
Valid until: 2024-05-26

Date, 2020-02-12

Christoph Dicks
Head of Certification/Notified Body

TÜV SÜD Product Service GmbH is Notified Body with identification no. 0123

TÜV SÜD Product Service GmbH • Certification Body • Ridlerstraße 65 • 80339 Munich • Germany

EC Certificate

Full Quality Assurance System
Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 039709 1263 Rev. 00

Facility(ies):

Medtronic Mexico S.de R.L.de CV
Av. Paseo Cucapah, 10510 El Lago, C.P. 22210 Tijuana, Baja California, MEXICO

Medtronic Perfusion Systems
7611 Northland Drive, Minneapolis, MN 55428, USA

Medtronic, Inc.
710 Medtronic Parkway, Minneapolis MN 55432, USA

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