



**CERTIFICAT**  
**privind lipsa sau existența restanțelor față de bugetul public național**

Nr.   
№

din   
от

**1. Destinația / Назначение**

Pentru participarea la proceduri de achiziții publice

**2. Date despre contribuabil / Информация о налогоплательщике**

Denumirea Наименование	Codul fiscal / Numărul de identificare Фискальный код / Идентификационный номер
<input type="text" value="BIOSISTEM MLD S.R.L."/>	<input type="text" value="1010600028048"/>
Adresa sediului de bază (strada, numărul) Адрес основного месторасположения (улица, номер)	Codul - Denumirea localității Код - Наименование населенного пункта
<input type="text" value="Albisoara nr.16 bl.1 of.7"/>	<input type="text" value="0150-SEC.RISCANI"/>

**3. Atestarea lipsei sau existenței restanțelor conform datelor Sistemului Informațional Automatizat /**

Подтверждение отсутствия или наличия недоимки согласно данных Информационной автоматизированной системы

La data emiterii prezentului certificat restanța față de bugetul public național constituie/ На дату выдачи данной справки недоимка перед национальным публичным бюджетом составляет:  
**0,00 lei/лей.**

**4. Valabil pînă la / Действителен до 19.01.2022**

**5. Autentificarea Serviciului Fiscal de Stat / Подтверждение Государственной налоговой службы**



L.S/ M.П.

Executor:

**Claudia GOJAN**  
Numele și prenumele/Fамилия и имя  
Tel.(022)323102

Semnătura/Подпись

**Petru GRICIUC**

Numele și prenumele/Фамилия и имя

Este extras din Sistemul Informațional al SFS SIA „Contul curent al contribuabilului”// 04.01.2022 ora 14:01:02  
cu aplicarea prevederilor pct. 82-83 Ordin IFPS nr.400 din 14.03.2014 (Monitorul Oficial 72-77/399, 28.03.2014)

NOTA (3,52)



Venituri financiare, total	090	490609	519239
din care:			
venituri din interese de participare	091		
inclusiv: veniturile obținute de la părțile afiliate	092		
venituri din dobânzi	093		25612
inclusiv: veniturile obținute de la părțile afiliate	094		
venituri din alte investiții financiare pe termen lung	095		
inclusiv: veniturile obținute de la părțile afiliate	096		
venituri aferente ajustărilor de valoare privind investițiile financiare pe termen lung și curente	097		
venituri din ieșirea investițiilor financiare	098		
venituri aferente diferențelor de curs valutar și de sumă	099	490609	493627
Cheltuieli financiare, total	100	686605	597528
din care:			
cheltuieli privind dobânzile	101		
inclusiv: cheltuielile aferente părților afiliate	102		
cheltuieli aferente ajustărilor de valoare privind investițiile financiare pe termen lung și curente	103		
cheltuieli aferente ieșirii investițiilor financiare	104		
cheltuieli aferente diferențelor de curs valutar și de sumă	105	686605	597528
<b>Rezultatul: profit (pierdere) financiar(ă)</b> (rd.090 - rd.100)	110	-195996	-78289
Venituri cu active immobilizate și excepționale	120		
Cheltuieli cu active immobilizate și excepționale	130		
<b>Rezultatul din operațiuni cu active immobilizate și excepționale: profit (pierdere)</b> (rd.120 - rd.130)	140		
<b>Rezultatul din alte activități: profit (pierdere)</b> (rd.110 + rd.140)	150	-195996	-78289
<b>Profit (pierdere) până la impozitare</b> (rd.080 + rd.150)	160	10081409	9025990
Cheltuieli privind impozitul pe venit	170	1178993	1051159
<b>Profit net (pierdere netă) al perioadei de gestiune</b> (rd.160 - rd.170)	180	8902416	7974831

**SITUAȚIA MODIFICĂRILOR CAPITALULUI PROPRIU**  
de la până la

Anexa 3

Nr. d/o	Indicatori	Cod rd	Sold la începutul perioadei de gestiune	Majorări	Diminuări	Sold la sfârșitul perioadei de gestiune
1	2	3	4	5	6	7
	<b>Capital social și neînregistrat</b>					
	1. Capital social	010				
	2. Capital nevărsat	020	( )	( )	( )	( )
	3. Capital neînregistrat	030				
I.	4. Capital retras	040	( )	( )	( )	( )
	5. Patrimoniul primit de la stat cu drept de proprietate	050				
	<b>Total capital social și neînregistrat</b> (rd.010 + rd.020 + rd.030 + rd.040 + rd.050)	060				
II.	<b>Prime de capital</b>	070				
	<b>Rezerve</b>					
	1. Capital de rezervă	080				
III.	2. Rezerve statutare	090				
	3. Alte rezerve	100				
	<b>Total rezerve</b> (rd.080 + rd.090 + rd.100)	110				
	<b>Profit (pierdere)</b>					
	1. Corecții ale rezultatelor anilor precedenți	120	X			

Версия для печати  
Сохранить

**Расписка 2**

Респондент  
Фискальный код: 1010600028048, наименование: BIOSISTEM MLD S.R.L.  
Предоставил отчет: RSE1\_21  
На фискальный период: A/2020  
Дата предоставления: 11.05.2021  
Временная метка отчета зарегистрированного в Информационной Системе НБС : 11.05.2021 12:26:31

National Bureau of Statistics (NBS) received the electronic version of the report, sent by you. The data provided is verified by NBS.

IV.	2. Profit nerepartizat (pierdere neacoperită) al anilor precedenți	130			
	3. Profit net (pierdere netă) al perioadei de gestiune	140	X		
	4. Profit utilizat al perioadei de gestiune	150	X	( )	( )
	<b>Total profit (pierdere)</b> (rd.120 + rd.130 + rd.140 + rd.150)	160			
V.	<b>Rezerve din reevaluare</b>	170			
VI.	<b>Alte elemente de capital propriu</b>	180			
	<b>Total capital propriu</b> (rd.060 + rd.070 + rd.110 + rd.160 + rd.170 + rd.180)	190			

**SITUAȚIA FLUXURILOR DE NUMERAR**  
de la până la

Anexa 4

Indicatori	Cod rd	Perioada de gestiune	
		precedentă	curentă
1	2	3	4
<b>Fluxuri de numerar din activitatea operațională</b>			
Încasări din vânzări	010		
Plăți pentru stocuri și servicii procurate	020		
Plăți către angajați și organe de asigurare socială și medicală	030		
Dobânzi plătite	040		
Plata impozitului pe venit	050		
Alte încasări	060		
Alte plăți	070		
<b>Fluxul net de numerar din activitatea operațională</b> (rd.010 - rd.020 - rd.030 - rd.040 - rd.050 + rd.060 - rd.070)	080		
<b>Fluxuri de numerar din activitatea de investiții</b>			
Încasări din vânzarea activelor immobilizate	090		
Plăți aferente intrărilor de active immobilizate	100		
Dobânzi încasate	110		
Dividende încasate	120		
inclusiv: dividende încasate din străinătate	121		
Alte încasări (plăți)	130		
<b>Fluxul net de numerar din activitatea de investiții</b> (rd.090 - rd.100 + rd.110 + rd.120 ± rd.130)	140		
<b>Fluxuri de numerar din activitatea financiară</b>			
Încasări sub formă de credite și împrumuturi	150		
Plăți aferente rambursării creditelor și împrumuturilor	160		
Dividende plătite	170		
inclusiv: dividende plătite rezidenților	171		
Încasări din operațiuni de capital	180		
Alte încasări (plăți)	190		
<b>Fluxul net de numerar din activitatea financiară</b> (rd.150 - rd.160 - rd.170 + rd.180 ± rd.190)	200		
<b>Fluxul net de numerar total</b> (± rd.080 ± rd.140 ± rd.200)	210		
Diferențe de curs valutar favorabile (nefavorabile)	220		
<b>Sold de numerar la începutul perioadei de gestiune</b>	230		
<b>Sold de numerar la sfârșitul perioadei de gestiune</b> (± rd.210 ± rd.220 + rd.230)	240		

**Documente atașate - Notă explicativă (fișierul pdf)**

Версия для печати  
Сохранить

**Расписка**

Респондент  
Фискальный код: 1010600028048, наименование: BIOSISTEM MLD S.R.L.  
Предоставил отчет: RSE1\_21  
На фискальный период: A/2020  
Дата предоставления: 11.05.2021  
Временная метка отчета зарегистрированного в Системе Электронной Отчетности и отправленного в Информационную Систему БНС : 11.05.2021 10:00:47

# SONY

Sony Belgium, bijkantoor van Sony Europe Limited  
Da Vincilaan 7 – D1, B-1935 Zaventem  
Phone: +32 (0) 2 706 43 11 - Fax : +32 (0) 2 706 43 20

---

## EU DECLARATION OF CONFORMITY

**Model Name: UPP-110S**

**93/42/EEC, 2007/47/EC, MDD**

# SONY

Sony Belgium, bijkantoor van Sony Europe Limited  
Da Vincilaan 7 – D1, B-1935 Zaventem  
Phone: +32 (0) 2 706 43 11 - Fax: +32 (0) 2 706 43 20

---

## EU DECLARATION OF CONFORMITY

1. Model No.:

**UPP-110S**

---

2. Name and address of the manufacturer's authorised representative:

**Sony Belgium, bijkantoor van Sony Europe Limited, Da Vincilaan 7-D1, 1935 Zaventem, Belgium**

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3. This declaration of conformity is issued under the sole responsibility of the manufacturer:

**Sony Corporation, 1-7-1 Konan Minato-ku Tokyo, 108-0075 Japan**

---

4. Object of the declaration:

**Thermal Print Media**

---

5. The object of the declaration described above is in conformity with:

**93/42/EEC, 2007/47/EC, MDD**

---

6. Where applicable, references to the relevant harmonised standards used or references to the technical specifications in relation to which conformity is declared:

**EN 60601-1:2006 + A1:2013**

---

7. Additional information:

**Following the provisions for Class I devices**

---

Signed for and on behalf of: **Sony Belgium, bijkantoor van Sony Europe Limited**

---

Zaventem, 2017-08-19



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Kris De Pauw  
Director  
Branch Manager

# SONY

Sony Belgium, bijkantoor van Sony Europe Limited  
Da Vinciilaan 7 – D1, B-1935 Zaventem  
Phone: +32 (0) 2 706 43 11 – Fax : +32 (0) 2 706 43 20

---

## EU DECLARATION OF CONFORMITY

**Model Name: UPP-110HG**

**93/42/EEC, 2007/47/EC, MDD**

# SONY

Sony Belgium, bijkantoor van Sony Europe Limited  
Da Vincilaan 7 – D1, B-1935 Zaventem  
Phone: +32 (0) 2 706 43 11 - Fax : +32 (0) 2 706 43 20

---

## EU DECLARATION OF CONFORMITY

1. Model No.:

**UPP-110HG**

---

2. Name and address of the manufacturer's authorised representative:

**Sony Belgium, bijkantoor van Sony Europe Limited, Da Vincilaan 7-D1, 1935 Zaventem, Belgium**

---

3. This declaration of conformity is issued under the sole responsibility of the manufacturer:

**Sony Corporation, 1-7-1 Konan Minato-ku Tokyo, 108-0075 Japan**

---

4. Object of the declaration:

**Thermal Print Media**

---

5. The object of the declaration described above is in conformity with:

**93/42/EEC, 2007/47/EC, MDD**

---

6. Where applicable, references to the relevant harmonised standards used or references to the technical specifications in relation to which conformity is declared:

**EN 60601-1:2006 + A1:2013**

---

7. Additional information:

**Following the provisions for Class I devices**

---

Signed for and on behalf of: **Sony Belgium, bijkantoor van Sony Europe Limited**

---

Zaventem, 2017-08-19



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Kris De Pauw  
Director  
Branch Manager



# SONY

Sony Belgium, bijkantoor van Sony Europe Limited  
Da Vinciiaan 7 – D1, B-1935 Zaventem  
Phone: +32 (0) 2 706 43 11 - Fax : +32 (0) 2 706 43 20

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## EU DECLARATION OF CONFORMITY

**Model Name: UPP-110HD**

93/42/EEC, 2007/47/EC, MDD

# SONY

Sony Belgium, bijkantoor van Sony Europe Limited  
Da Vincilaan 7 – D1, B-1935 Zaventem  
Phone: +32 (0) 2 706 43 11 - Fax : +32 (0) 2 706 43 20

---

## EU DECLARATION OF CONFORMITY

1. Model No.:  
**UPP-110HD**

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2. Name and address of the manufacturer's authorised representative:  
**Sony Belgium, bijkantoor van Sony Europe Limited, Da Vincilaan 7-D1, 1935 Zaventem, Belgium**

---

3. This declaration of conformity is issued under the sole responsibility of the manufacturer:  
**Sony Corporation, 1-7-1 Konan Minato-ku Tokyo, 108-0075 Japan**

---

4. Object of the declaration:  
**Thermal Print Media**

---

5. The object of the declaration described above is in conformity with:  
**93/42/EEC, 2007/47/EC, MDD**

---

6. Where applicable, references to the relevant harmonised standards used or references to the technical specifications in relation to which conformity is declared:  
**EN 60601-1:2006 + A1:2013**

---

7. Additional information:  
**Following the provisions for Class I devices**

---

Signed for and on behalf of: **Sony Belgium, bijkantoor van Sony Europe Limited**

---

Zaventem, 2017-08-19



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Kris De Pauw  
Director  
Branch Manager

# EC Certificate

## Full Quality Assurance System

Certificate No.:  
**245506-2017-CE-IND-NA-PS Rev. 2.0**

Project No.:  
**PRJC-499089-2014-MSL-IND**

Valid Until:  
**27 May 2024**

This is to certify that the quality system of:

### **Meril Endo Surgery Pvt Ltd**

Third Floor, E1- E3, Meril Park, Survey No 135/2/B & 174/2, Muktanand Marg,  
Chala, Vapi, Gujarat, India - 396191

For design, production and final product inspection/testing of:

### **Sterile / non - sterile surgical sutures with and without needle**

Has been assessed with respect to:

### **The conformity assessment procedure described in Annex II of Council Directive 93/42/EEC on Medical Devices, as amended**

and found to comply.

Further details of the product(s) and conditions for certification are given overleaf.

Place and Date:  
**Høvik, 16 July 2019**



For:  
**DNV GL PRESAFE AS**



**Cathrine Wisbech**

The Certificate has been digitally signed.  
See [www.presafe.com/digital\\_signatures](http://www.presafe.com/digital_signatures) for more info

Notice: The Certificate is subject to terms and conditions as set out in the Certification Agreement. Failure to comply may render this Certificate invalid.

# EC Certificate

## Full Quality Assurance System

Certificate No.:  
**245506-2017-CE-IND-NA-PS Rev. 2.0**

Project No.:  
**PRJC-499089-2014-MSL-IND**

Valid Until:  
**27 May 2024**

### Jurisdiction

Application of Council Directive 93/42/EEC of 14 June 1993, adopted as “Forskrift om Medisinsk Utstyr” by the Norwegian Ministry of Health and Care Services.

### Certificate history:

Revision	Description	Issue Date
0.0	Supersedes DNVGL (NB 0434) Certificate no: 151561-2014-CE-IND-NA Rev 3.0 following transfer to notified body functions to DNV GL Nemko Presafe AS (NB 2460)	2017-11-15
1.0	Remove of Polypropylene Mesh	2018-01-08
2.0	<b>Recertification and reduction in scope</b>	2019-07-16

### Products covered by this Certificate:

Product Description	Product Name	Class
Absorbable Sutures	<ul style="list-style-type: none"> <li>Megasorb™ / Aspiron™ Polyglycolic acid Braided coated Polyglycolic acid suture</li> <li>Mitsu™ / Aspiron™ Polyglactin 910 and Mitsu FST™ / Aspiron™ Polyglactin 910 FST. Braided coated Poly (glycolide/l-lactide) suture</li> <li>Filaxyn™ / Aspiron™ Polydioxanone suture. Monofilament Poly (p-dioxanone) suture</li> <li>Filapron™ / Aspiron™ Polyglecaprone 25 suture. Monofilament poly (glycolide-co-caprolactone) suture</li> </ul>	III*
Non-Absorbable Sutures	<ul style="list-style-type: none"> <li>Filaprop™ / Aspiron™ Polypropylene Blue Monofilament Polypropylene Suture</li> </ul>	III**



# EC Certificate

## Full Quality Assurance System

Certificate No.:  
**245506-2017-CE-IND-NA-PS Rev. 2.0**

Project No.:  
**PRJC-499089-2014-MSL-IND**

Valid Until:  
**27 May 2024**

\*Design assessment is covered by a separate EC-Design Examination Certificate No.: 245507-2017-CE-IND-NA-PS Rev. 2

\*\*Design assessment is covered by a separate EC-Design Examination Certificate No.: 245508-2017-CE-IND-NA-PS Rev. 2

### Sites covered by this certificate

Site Name	Address
Meril Endo Surgery Pvt Ltd	Third Floor, E1- E3, Meril Park, Survey No 135/2/B & 174/2, Muktanand Marg, Chala, Vapi, Gujarat, India - 396191

### EU Representative

OBELIS S.A

Bd. Général Wahis, 53, 1030 Brussels, Belgium. Tel: +32.2.732.59.54. Fax: +32.2.732.60.03

E-mail: [mail@obelis.net](mailto:mail@obelis.net), [www.obelis.net](http://www.obelis.net)

# EC Certificate

## Full Quality Assurance System

Certificate No.:  
**245506-2017-CE-IND-NA-PS Rev. 2.0**

Project No.:  
**PRJC-499089-2014-MSL-IND**

Valid Until:  
**27 May 2024**

### Terms and conditions

The certificate is subject to the following terms and conditions:

- Any producer (see 2001/95/EC for a precise definition) is liable for damage caused by a defect in his product(s), in accordance with directive 85/374/EEC, as amended, concerning liability of defective products.
- The certificate is only valid for the products and/or manufacturing premises listed above.
- The Manufacturer shall fulfil the obligations arising out of the quality system as approved and uphold it so that it remains adequate and efficient.
- The Manufacturer shall inform Presafe of any intended updating of the quality system and Presafe will assess the changes and decide if the certificate remains valid.
- Periodical audits will be held, in order to verify that the Manufacturer maintains and applies the quality system. Presafe reserves the right, on a spot basis or based on suspicion, to pay unannounced visits.

The following may render this Certificate invalid:

- Changes in the quality system affecting production.
- Periodical audits not held within the allowed time window.

### Conformity declaration and marking of product

When meeting with the terms and conditions above, the producer may draw up an EC declaration of conformity and legally affix the CE mark followed by the Notified Body identification number of Presafe.

End of Certificate



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 077790 0060 Rev. 00**

**Manufacturer:**

**Covidien LLC**

15 Hampshire Street  
Mansfield MA 02048  
USA

**Product Category(ies): Oximetry and Capnography Monitor Systems  
Temperature Monitor Systems, Patient Warming  
Device Systems, Disposable Airway Management  
Devices, Tracheal Tubes, Tracheostomy Tubes,  
Speaking Valves, and Intubating Stylets, Ventilator  
Systems and Patient Interface Circuit Systems,  
EEG Monitoring Systems, Breathing Therapy and  
Humidification, Heated Inspiratory Line  
Humidifiers, Multi-patient Physiologic Monitoring  
System and Data Analytics Software,  
Gastrointestinal Measurement and Dilation System,  
Electrosurgical Diathermy System Electrode.**

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 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.:** 72145607

**Valid from:** 2020-06-29

**Valid until:** 2024-05-26

**Date,** 2020-06-29

Christoph Dicks  
Head of Certification/Notified Body