

EC Certificate Full Quality Assurance System: Certificate KR10/01512

The management system of

## 3D Imaging & Simulations Corp.

815, Tamnip-Dong, Yuseong-Gu, Daejeon, Korea

has been assessed and certified as meeting the requirements of

### Directive 93/42/EEC

on medical devices, Annex II (excluding Section 4)

For the following products

**Computed Radiography Scanner system. (Model: FireCR, FireCR Dental, Dental CR, Planmeca ProScanner)**

For placing on the market of Class III devices covered by this certificate, an EC Design Examination Certificate according to Annex II (Section 4) is required.

This certificate is valid from 8 October 2013 until 8 October 2018 and remains valid subject to satisfactory surveillance audits.

Re certification audit due before 29 September 2016

Issue 6. Certified since 8 October 2010

Certification is based on reports numbered KR/SEL Y-PC/10245

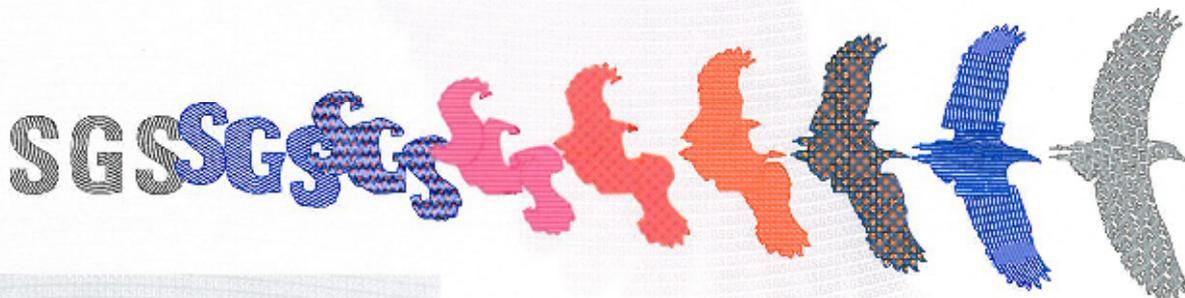
Authorised by

**SGS United Kingdom Ltd, Notified Body 0120**

202B Worle Parkway, Weston-super-Mare, BS22 6WA UK  
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SGS CE 01 0311

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EB Sertifikatas Pilna kokybės užtikrinimo sistema: Sertifikatas KR10/01512

Vadybos sistema

**3D Imaging & Simulations Corp.**  
815, Tamnip-Dong, Yuseong-Gu, Daejeon, Korėja

įvertinto ir sertifikavo kaip atitinkanti

Medicinos prietaisų, II Priedo (išskyrus 4 dalį)

**93/42/EEB direktyvos**  
reikalavimus

Šių produktų

Skaitmeninė rentgenografijos nuskaitymo sistema. (Modelis: FireCR, ForeCR  
Dental, Dental CR, Planmeca ProScanner)

Įvedant į rinką III klasės prietaisus, kuriems taikomas šis sertifikatas,  
pagal II priedo (4 dalį) reikalingas EB konstrukcijos tyrimo sertifikatas.

Šis sertifikatas galioja nuo 2013 m. spalio 8 d. iki 2018 m. spalio 8 d.  
Lieka galioti priklausomai nuo patenkinamų priežiūros audito rezultatų.  
Pakartotinis atestavimas turi būti atliktas prieš 2016 m. rugsėjo 29 d.  
6 leidimas. Sertifikuota nuo 2010 m. spalio 8 d.

Šis Sertifikatas remiasi sprendimu: KR/SEL Y-PC/10245

SGS United Kingdom Ltd., Patvirtinanti įstaiga 0120  
202B Worle Parkway, Weston-super-Mare, BS22 6WA UK

SGS CE 01 0311



Vertimas teisingas  
UAB „Plandent“ direktorius Andrius Preišegalavičius





# EC-CERTIFICATE

(Full quality assurance system)



This is to certify that the company

**KLS martin**  
GROUP

## Gebrüder Martin GmbH & Co. KG

KLS Martin Platz 1  
78532 Tuttlingen  
Germany

has implemented and maintains a full quality assurance system which applies to the products at every stage from design to final controls.

Through an audit, documented in a report, performed by DQS Medizinprodukte GmbH, it was verified that the management system fulfills the requirements of

### Annex II – excluding Section 4 of Council Directive 93/42/EEC concerning medical devices

with respect to the following medical devices:

Active and non active medical devices; surgical instruments and units, implants, medical devices for RF- and laser surgery, HF- and laser surgery, ceiling-mounted supply systems and accessories according to annex.

The manufacturer is subject to surveillance according to Annex II, Section 5. The CE marking with the Notified Body Identification Number (0297) may be affixed on the devices listed in the certificate. An EC Design Examination Certificate according to Annex II, Section 4 is required for class III devices covered by this certificate. The certificate is in the case of class I(s) devices (I(s) = class I products placed on the market in sterile conditions) limited to the aspects of manufacture concerned with securing and maintaining sterile conditions. The certificate is in the case of class I(m) devices (I(m) = class I devices with a measuring function) limited to the aspects of manufacture concerned with the conformity of the products with the metrological requirements.

Certificate registration no.	210299 MR2
Certificate unique ID	170639389
Effective date	2016-01-04
Expiry date	2019-05-26
Frankfurt am Main	2016-01-04

### DQS Medizinprodukte GmbH

Frank Graichen  
Managing Director

Dr. Thomas Feldmann  
Head of Certification Body

August-Schanz-Straße 21, 60433 Frankfurt am Main,  
Tel. +49 (0) 69 95427-300, [medical.devices@dqs-med.de](mailto:medical.devices@dqs-med.de)

DQS Medizinprodukte GmbH is a Notified Body according to Council Directive 93/42/EEC concerning medical devices with the Identification Number 0297.



**Annex to Certificate**  
**Certificate registration No.: 210299 MR2**  
**Certificate unique ID: 170639389**  
**Effective date: 2016-01-04**



## **Gebrüder Martin GmbH & Co. KG**

KLS Martin Platz 1  
78532 Tuttlingen  
Germany

<b>Device family</b>	<b>Devices</b>	<b>Class</b>
RF-surgery units	Medical RF-surgery and HF-surgery units	IIb
HF-surgery units	for general and special applications Argon Beamer	IIb
Accessories for RF-surgery units	Handles for RF-surgery and HF-surgery	IIa / IIb
Accessories for HF-surgery units	Electrodes Forceps and scissors Clamps, forceps, NervScout	
Laser surgery units	surgical laser units for common and particular use	IIb
Accessories for laser surgery units	Laser contact probe Adapter for laser fibres Connection fibres Fibre optics Bipolar suction and irrigation units Micromanipulator	IIb
Smoke evacuation systems	Accessories for Smoke evacuation systems	IIa
Accessories for Smoke evacuation systems	Adapter	IIa
Implants and Instruments	Distractor Trocar tip for activation arm Condylar implant Washers Meshes (for cranioplasty) Bone plates Bone screws Bone nails Fixation screws Fixation screws (lug) UHP (prosthesis for wrist joint) Tantal wire (orthodontic) Bone clips Bone wire Sternal Talon Finger joint	IIa / IIb
	Aneurysm Clip system permanent and temporary	III



**Annex to Certificate**  
**Certificate registration No.: 210299 MR2**  
**Certificate unique ID: 170639389**  
**Effective date: 2016-01-04**



## **Gebrüder Martin GmbH & Co. KG**

KLS Martin Platz 1  
78532 Tuttlingen  
Germany

<b>Device family</b>	<b>Devices</b>	<b>Class</b>
Resorbable Implants	Resorb X meshes Resorb X plates Resorb X screws Resorb X Pins	III
Surgical instruments and units	Sterile scalpels Sterile blades Guide pins Guide wires  Drill bits OMF surgery BOS driver SonicWeld Rx (basis device, handpieces) SonicWeld Rx (sonotrodes) Drill bits / Saw blades orthopaedic BOS-Battery pack sterile Bending Templates	IIa IIa IIb IIb  IIa IIa IIa IIa IIa Is Is
Pendants	Service Head	IIb
Bone Grafting Material	Martrix	III

Vertimas iš anglų kalbos į lietuvių kalbą

CE – Sertifikatas  
(Pilna kokybės užtikrinimo sistema)

Skirtas patvirtinti, kad kompanija  
KLS Martin Group  
Gebrüder Martin GmbH & CO. KG  
KLS Martin Platz 1  
78532 Tuttlingen  
Vokietija

Įdiegė ir palaiko pilną kokybės valdymo sistemą, kuri taikoma produktams kiekviename etape nuo jų kūrimo iki galutinės kontrolės.

DQS Medizinprodukte GmbH atliktas auditas, dokumentuotas ataskaita, patvirtino, kad ši kokybės valdymo sistema atitinka

93/42/EEC medicininių įrenginių Direktyvos, II Priedo – išskyrus 4 Skyrių , reikalavimus

ir pritaikyta šiems medicininiams įrenginiams:

Veikiantys ir neveikiantys medicininiai įrenginiai; chirurgijos instrumentai ir elementai, implantai, medicininiai įrenginiai RF- ir lazerio chirurgijai, HF- ir lazerinei chirurgijai, lubinės sistemos ir aksesuarai pagal priedą

Kaip numatyta minėtos direktyvos II Priedo 5 skyriuje, gamintojas yra periodinio tikrinimo subjektas. CE ženklas su įgaliosios įstaigos identifikacijos numeriu (0297) gali būti naudojamas su sertifikate išvardintais gaminiais. CE Konstruktijos Patikrinimo Sertifikatas pagal II Priedo, 4 Skyriaus reikalavimus yra reikalingas šiame sertifikate išvardintiems III klasės įrenginiams. Sertifikatas yra ribojamas I(s) klasės įrenginiams (I(s) = I klasės produktai, tiekiami į rinką steriliomis sąlygomis) atsižvelgiant į gamintojo suinteresuotumą užtikrinant ir palaikant sterilias sąlygas. Sertifikatas yra ribojamas I(m) klasės įrenginiams (I(m) = I klasės produktai su matavimo funkcija) atsižvelgiant į gamintojo suinteresuotumą užtikrinant prietaisų atitikimą metrologiniams reikalavimams.

Sertifikato registracijos Nr.	210299 MR2
Sertifikato unikalus ID	170639389
Įsigaliojimo data	2016-01-04
Galioja iki	2019-05-26
Frankfurtas prie Maino	2016-01-04

DQS Medizinprodukte GmbH

/parašas/

Frank Graichen  
Generalinis direktorius

/parašas/

Dr. Thomas Feldmann  
Sertifikavimo įstaigos vadovas

August-Schanz-Strasse 21, 60433 Frankfurt am Main,

Tel. +49 (0) 69 95427-300, [medical.devices@dqs.de](mailto:medical.devices@dqs.de)

DQS Medizinprodukte GmbH yra pagal Tarybos Direktyvą 93/42/EEC, susijusią su mediciniais įrenginiais, Įgaliosas Asmuo su Identifikacijos numeriu: 0297

Priedas prie sertifikato  
 Sertifikato registracijos Nr.: 210299 MR2  
 Sertifikato unikalus ID: 170639389  
 Įsigaliojimo data: 2016-01-04

Gebrüder Martin GmbH & Co. KG  
 KLS Martin Platz 1  
 78532 Tuttlingen  
 Vokietija

Produktų grupės	Produktai	Klasė
RF- chirurgijos įrenginiai HF- chirurgijos įrenginiai	Medicininiai RF-chirurgijos ir HF-chirurgijos įrenginiai bendram ir specialiam vartojimui Argono spinduliavimo įrenginys	I <b>b</b> I <b>b</b>
RF- chirurgijos įrenginių priedai HF- chirurgijos įrenginių priedai	Laikikliai RF-chirurgijai ir HF-chirurgijai Elektrodai Pincetai ir žirklys Spaustukai, chirurginės žnyplės NervScout	I <b>a</b> / I <b>b</b>
Lazerinės chirurgijos įrenginiai	Chirurginiai lazeriniai įrenginiai bendram ir konkrečiam naudojimui	I <b>b</b>
Lazerinės chirurgijos įrenginių priedai	Lazerio kontaktinis zondas Lazerio pluoštų adapteris Pluoštų sąsaja Skaidulų optika Bipoliario siurbimo ir drėkinimo įrenginiai Mikromanipulatorius	I <b>b</b>
Dūmų evakuacijos sistemos	Priedai dūmų evakuacijos sistemai	I <b>a</b>
Priedai dūmų evakuacijos sistemai	Adapteris	I <b>a</b>
Implantai ir instrumentai	Distraktorius Trokaro antgalis aktyvacijos rankenai Krumplių implantas Kraujagyslių spaustukai Tinklelis Kaulo plokštelės Kaulo sraigtai Kaulo vinys Fiksavimo sraigrai Fiksavimo sraigtai (tempimas) UHP (riešo sąnario protezai) Tantalinė viela (ortodontinė) Kaulų sąvaržos Kaulų viela Krūtinkaulio fiksatorius Piršto sąnarys	I <b>a</b> / I <b>b</b>
	Laikina ir nuolatinė aneurizmos spaustukų sistema	I <b>II</b>

Priedas prie sertifikato  
Sertifikato registracijos Nr.: 210299 MR2  
Sertifikato unikalus ID: 170639389  
Įsigaliojimo data: 2016-01-04

Gebrüder Martin GmbH & Co. KG  
KLS Martin Platz 1  
78532 Tuttlingen  
Vokietija

Produktų grupės	Produktai	Klasė
Rezorbuojami implantai	Rezorbuojami X tinkleliai Rezorbuojamos X plokštės Rezorbuojami X sraigčiai Rezorbuojami X smeigtukai	III
Chirurginiai instrumentai ir įrenginiai	Sterilūs skalpeliai Sterilūs peiliukai Smeigtukai Vielos	IIa IIa IIb IIb
	OMF chirurgijos gražtai BOS suktuvas SonicWeld Rx (pagrindinis įrenginys) SonicWeld Rx (sonotrodas) Ortopediniai gražtai / pjūklai Boss-Battery pakuotė sterili	IIa IIa IIa IIa IIa Is
Kaulinė medžiaga	Matrix	III

Šis priedas galioja tik kartu su aukščiau paminėtu sertifikatu.

Vertimą tvirtino UAB „Graina“ direktorius valdymui Šarūnas Gildutis





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 14 01 45735 027

**Manufacturer:** **3M ESPE Dental Products**  
2510 Conway Avenue  
St. Paul MN 55144-1000  
USA

**EC-Representative:** **3M Deutschland GmbH**  
Carl-Schurz-Straße 1  
41453 Neuss  
GERMANY

**Product Category(ies):** **Sterile and Non-Sterile Dental Products including: Resin Based Restorative Materials, Dental Adhesives, Primers and Cements, Pit and Fissure Sealants, Glass Ionomers, Pre-Formed Crowns, Etchants, Cavity Varnishes and Liners, Dental and Orthodontic Endosseous Implants, Prosthetic Components, Orthodontic Springs, Dental and Surgical Instruments, Denture Relining Material**

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.:** NM1212860

**Valid from:** 2014-02-07

**Valid until:** 2016-12-03

Hans-Heiner Junker



**Date,** 2014-02-07

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



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 14 01 45735 027**

**Facility(ies):**

3M ESPE Dental Products  
2510 Conway Avenue, St. Paul MN 55144-1000, USA

3M ESPE Dental Products  
2111 McGaw Avenue, Irvine CA 92614, USA



Product Service

# EC SERTIFIKATAS

Pilnas kokybės sistemos patvirtinimas

Mediciniu prietaisų 93/42/EEB direktyva, II priedas, išskyrus (4)

(Prietaisai IIa, IIb ar III klasės)

**Nr. G1 14 01 45735 027**

**Gamintojas:**

**3M ESPE Dental Products**

3M Conway Avenue  
St. Paul, MN 55144 - 1000  
USA (JAV)

**EC-atstovas:**

**3M Deutschland GmbH**

Carl-Schurz-Staße 1,  
41453 Neuss,  
Germany (Vokietija)

**Priemonės(-ių)  
kategorija:**

**Sterilūs ir nesterilūs odontologiniai produktai įskaitant:  
dervinės restauracinės medžiagos, odontologiniai adhesyvai,  
praimeriai ir cementai, vagelių ir įtrūkimų silantai,  
stiklo jonomerai, karūnėlių pusfabrikačiai, ėsdintojai,  
ertmių lakai ir laineriai, odontologiniai ir ortodontiniai  
endoosaliniai implantai, protezavimo komponentai,  
ortodontinės spyruoklės, odontologiniai ir chirurginiai  
instrumentai, dantų protezų perbazavimo medžiagos**

TÜV SÜD Product Service GmbH, sertifikavimo institucija patvirtina Kokybės garantijos sistemą taikomą dizainui, gamybai ir galutiniam patikrinimui viršuje paminėtam gamintojui, atitinkamiems produktams / produktų kategorijoms pagal medicinos prietaisų 93/42/EEB direktyvos II priedo dalį. Ši kokybės valdymo sistema atitinka šios direktyvos reikalavimus ir yra periodiškai tikrinama. III klasės prietaisų rinkodarai yra privalomas papildomas II (4) priedo sertifikatas.

**Pranešimo Nr.:**

**NM1105459**

Galioja nuo:

2014-02-07

Galiojimo data:

2016-12-03

**Data: 2014-02-07**

**Hans-Heiner Junker**

VERTIMAS TEISINGAS  
UAB „Plandent“ direktorius Andrius Preišegalavičius

TÜV SÜD Product Service GmbH yra Sertifikavimo įstaiga, kurios identifikacijos Nr. 0123.

# EC SERTIFIKATAS

**Pilnas kokybės sistemos patvirtinimas**

**Medicinių prietaisų 93/42/EEB direktyva, II priedas, išskyrus (4)**

**(Prietaisai IIa, IIb ar III klasės)**

**Nr. G1 14 01 45735 027**



Product Service

**Įrenginys(iai):**

3M ESPE Dental Products

2510 Conway Avenue, St.Paul, MN 55144 – 1000, USA (JAV)

3M ESPE Dental Products

2111 McGaw Avenue, Irvine, CA 92614, USA (JAV)



**EC DECLARATION OF CONFORMITY**

Annex VII of Council Directive 93/42/EEC concerning Medical Devices.

**Manufacturer:**

3M ESPE Dental Products  
3M Center, Bldg. 260  
St. Paul, Minnesota 55144-1000  
USA

ISO Certificate No: G1 11 11 45735 020  
Valid until: 2016-12-03

GMDN CODE 35807 – Discs, dental abrasive  
35702 – Abrasive strips, dental  
35170 – Mandrel, dental

UMDNS CODE 16-201 – Polishing System  
11-282 – Grinding wheel, dental

**Manufacturing Facility:**

3M ESPE Dental Products  
2111 McGaw Avenue  
Irvine, California 92614  
USA

Product Family: Abrasives

- Product:
- Sof-Lex™ Contouring and Polishing Discs
  - Sof-Lex™ Finishing and Polishing Discs
  - Sof-Lex™ Finishing and Polishing Strips
  - Sof-Lex™ Extra Thin Contouring and Polishing Discs
  - Sof-Lex™ Finishing and Polishing Systems
  - Sof-Lex™ Mandrels

CE Mark Date: August 1994

EC Product Class: I  
Rule:

Per Annex IX of the Medical Device Directive

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**Declaration of Conformity**

3M ESPE Dental Products declares that devices on the attached Device Schedule conform to the relevant provisions of the EC Council Directive 93/42/EEC dated 14 June 1993 and is in accordance with ISO 13485:2003, Medical devices – Quality management systems – Requirements for regulatory purposes, as implemented by the European Union's Medical Devices Regulations.

3M ESPE Dental Products has appointed 3M Deutschland GmbH as our EU Authorized Representative.  
3M Deutschland GmbH  
Carl-Schurz-Strasse 1, 41453 Neuss, Germany

Director - 3M ESPE  
Regulatory Affairs St. Paul, Minnesota USA 20 January 2012  
Title Location Date

**EC DECLARATION OF CONFORMITY**

**Product Family:**

Abrasives

**Device Description:**

See below

**EC Product Class (Per Annex IX Rules):**

Class: I Rule: 5 of Annex IX of the Medical Device Directive

- Sof-Lex™ Contouring and Polishing Discs
- Sof-Lex™ Finishing and Polishing Discs
- Sof-Lex™ Finishing and Polishing Strips
- Sof-Lex™ Extra Thin Contouring and Polishing Discs
- Sof-Lex™ Finishing and Polishing Systems
- Sof-Lex™ Mandrels

**Product:**

<b>Western Europe</b>	
<b>Catalog/ Order Number</b>	<b>Description</b>
	<b>Sof-Lex™ Contouring and Polishing Discs</b>
1980	Sof-Lex Contouring and Polishing Disc Kit <ul style="list-style-type: none"> <li>• 30 – 1/2 inch (12,7mm) Discs – Coarse</li> <li>• 30 – 1/2 inch (12,7mm) Discs – Medium</li> <li>• 30 – 1/2 inch (12,7mm) Discs – Fine</li> <li>• 30 – 1/2 inch (12,7mm) Discs – Super Fine</li> <li>• 30 – 3/8 inch (9,5mm) Discs – Coarse</li> <li>• 30 – 3/8 inch (9,5mm) Discs – Medium</li> <li>• 30 – 3/8 inch (9,5mm) Discs – Fine</li> <li>• 30 – 3/8 inch (9,5mm) Discs – Super Fine</li> <li>• 1 – RA Mandrel</li> </ul>
1981 C	85 – 3/8 inch (9,5mm) Sof-Lex Contouring and Polishing Discs - Coarse
1981 M	85 – 3/8 inch (9,5mm) Sof-Lex Contouring and Polishing Discs - Medium
1981 F	85 – 3/8 inch (9,5mm) Sof-Lex Contouring and Polishing Discs - Fine
1981 SF	85 – 3/8 inch (9,5mm) Sof-Lex Contouring and Polishing Discs – Super Fine
1982 C	85 – 1/2 inch (12,7mm) Sof-Lex Contouring and Polishing Discs - Coarse
1982 M	85 – 1/2 inch (12,7mm) Sof-Lex Contouring and Polishing Discs - Medium
1982 F	85 – 1/2 inch (12,7mm) Sof-Lex Contouring and Polishing Discs - Fine
1982 SF	85 – 1/2 inch (12,7mm) Sof-Lex Contouring and Polishing Discs – Super Fine
1981 GE	Sof-Lex Studentenset (German Version)
	<b>Sof-Lex™ Finishing and Polishing Discs</b>
1958 C	100 – 5/8 inch Square Eyelet Discs - Coarse
1958 M	100 – 5/8 inch Square Eyelet Discs - Medium
1958 F	100 – 5/8 inch Square Eyelet Discs - Fine
1958 SF	100 – 5/8 inch Square Eyelet Discs – Super Fine
	<b>Sof-Lex™ Finishing and Polishing Strips</b>
1956	120 – 7 inches x 5/32 inch (178mm x 4mm) Sof-Lex Finishing and Polishing Strips – Fine/Super Fine
1954	150 – 7 inches x 5/32 inch (178mm x 4mm) Sof-Lex Finishing and Polishing Strips – Coarse/Medium
1954 N	100 - 7 inches x 5/64 inch (178mm x 2mm) Sof-Lex Finishing and Polishing Strips – Coarse/Medium, Narrow

<b>Sof-Lex™ Extra Thin Contouring and Polishing Discs</b>	
2380	<p>Sof-Lex Extra Thin (XT) Contouring and Polishing Disc Kit</p> <ul style="list-style-type: none"> <li>• 30 – 1/2 inch (12,7mm) XT Discs – Coarse</li> <li>• 30 – 1/2 inch (12,7mm) XT Discs – Medium</li> <li>• 30 – 1/2 inch (12,7mm) XT Discs – Fine</li> <li>• 30 – 1/2 inch (12,7mm) XT Discs – Super Fine</li> <li>• 30 – 3/8 inch (9,5mm) XT Discs – Coarse</li> <li>• 30 – 3/8 inch (9,5mm) XT Discs – Medium</li> <li>• 30 – 3/8 inch (9,5mm) XT Discs – Fine</li> <li>• 30 – 3/8 inch (9,5mm) XT Discs – Super Fine</li> <li>• 1 – RA Mandrel</li> </ul>
2381 C	85 – 3/8 inch (9,5mm) Sof-Lex Extra Thin (XT) Contouring and Polishing Discs - Coarse
2381 M	85 – 3/8 inch (9,5mm) Sof-Lex Extra Thin (XT) Contouring and Polishing Discs - Medium
2381 F	85 – 3/8 inch (9,5mm) Sof-Lex Extra Thin (XT) Contouring and Polishing Discs - Fine
2381 SF	85 – 3/8 inch (9,5mm) Sof-Lex Extra Thin (XT) Contouring and Polishing Discs – Super Fine
2382 C	85 – 1/2 inch (12,7mm) Sof-Lex Extra Thin (XT) Contouring and Polishing Discs - Coarse
2382 M	85 – 1/2 inch (12,7mm) Sof-Lex Extra Thin (XT) Contouring and Polishing Discs - Medium
2382 F	85 – 1/2 inch (12,7mm) Sof-Lex Extra Thin (XT) Contouring and Polishing Discs - Fine
2382 SF	85 – 1/2 inch (12,7mm) Sof-Lex Extra Thin (XT) Contouring and Polishing Discs – Super Fine
<b>Sof-Lex™ Finishing and Polishing Systems Kit</b>	
2385 P	<p>Sof-Lex Finishing and Polishing Systems Kit</p> <ul style="list-style-type: none"> <li>• 30 – 1/2 inch (12,7mm) Discs – Coarse</li> <li>• 30 – 1/2 inch (12,7mm) Discs – Medium</li> <li>• 30 – 1/2 inch (12,7mm) Discs – Fine</li> <li>• 30 – 1/2 inch (12,7mm) Discs – Super Fine</li> <li>• 30 – 3/8 inch (9,5mm) Discs – Coarse</li> <li>• 30 – 3/8 inch (9,5mm) Discs – Medium</li> <li>• 30 – 3/8 inch (9,5mm) Discs – Fine</li> <li>• 30 – 3/8 inch (9,5mm) Discs – Super Fine</li> <li>• 30 – 1/2 inch (12,7mm) XT Discs – Coarse</li> <li>• 30 – 1/2 inch (12,7mm) XT Discs – Medium</li> <li>• 30 – 1/2 inch (12,7mm) XT Discs – Fine</li> <li>• 30 – 1/2 inch (12,7mm) XT Discs – Super Fine</li> <li>• 30 – 3/8 inch (9,5mm) XT Discs – Coarse</li> <li>• 30 – 3/8 inch (9,5mm) XT Discs – Medium</li> <li>• 30 – 3/8 inch (9,5mm) XT Discs – Fine</li> <li>• 30 – 3/8 inch (9,5mm) XT Discs – Super Fine</li> <li>• 15 – 7 inches x 5/32 inch (178mm x 4mm) Strips – Coarse/Medium</li> <li>• 15 – 7 inches x 5/64 inch (178mm x 2mm) Narrow Strips – Coarse/Medium</li> <li>• 15 – 7 inches x 5/32 inch (178mm x 4mm) Strips – Fine/Super Fine</li> <li>• 1 – RA Mandrel</li> </ul>
2385 MP	<p>Sof-Lex Finishing and Polishing System Sample Kit</p> <ul style="list-style-type: none"> <li>• 1- Sof-Lex Polishing Strips - Fine/Superfine</li> <li>• 1- Sof-Lex Polishing Strips - Coarse/Medium, Narrow</li> <li>• 1- Sof-Lex Polishing Strips - Coarse/Medium</li> </ul>
2385 GE	Sof-Lex Mustertute (German Version)
2385 ST	Sof-Lex Set Heidelberg (German Version)

<b>Sof-Lex™ Mandrels</b>	
1983 RA	Sof-Lex RA Mandrels (3)
1983 HP	Sof-Lex HP Mandrels (3)
1983 FG	Sof-Lex FG Mandrels (3)
<b>Eastern Europe</b>	
<b>Catalog/ Order Number</b>	<b>Description</b>
<b>Sof-Lex™ Contouring and Polishing Discs</b>	
8691C	50 – 1/2 inch (12,7mm) Sof-Lex Contouring and Polishing Discs - Coarse
8691F	50 – 1/2 inch (12,7mm) Sof-Lex Contouring and Polishing Discs - Medium
8691M	50 – 1/2 inch (12,7mm) Sof-Lex Contouring and Polishing Discs - Fine
8691SF	50 – 1/2 inch (12,7mm) Sof-Lex Contouring and Polishing Discs – Super Fine
8690C	50 – 3/8 inch (9,5mm) Sof-Lex Contouring and Polishing Discs - Coarse
8690M	50 – 3/8 inch (9,5mm) Sof-Lex Contouring and Polishing Discs - Medium
8690F	50 – 3/8 inch (9,5mm) Sof-Lex Contouring and Polishing Discs - Fine
8690SF	50 – 3/8 inch (9,5mm) Sof-Lex Contouring and Polishing Discs – Super Fine
<b>Sof-Lex™ Extra Thin Contouring and Polishing Discs</b>	
8692C	50 – 1/2 inch (12,7mm) Sof-Lex Extra Thin (XT) Contouring and Polishing Discs - Coarse
8692M	50 – 1/2 inch (12,7mm) Sof-Lex Extra Thin (XT) Contouring and Polishing Discs - Medium
8692F	50 – 1/2 inch (12,7mm) Sof-Lex Extra Thin (XT) Contouring and Polishing Discs - Fine
8692SF	50 – 1/2 inch (12,7mm) Sof-Lex Extra Thin (XT) Contouring and Polishing Discs – Super Fine
8693C	50 – 3/8 inch (9,5mm) Sof-Lex Extra Thin (XT) Contouring and Polishing Discs - Coarse
8693F	50 – 3/8 inch (9,5mm) Sof-Lex Extra Thin (XT) Contouring and Polishing Discs - Medium
8693M	50 – 3/8 inch (9,5mm) Sof-Lex Extra Thin (XT) Contouring and Polishing Discs - Fine
8693SF	50 – 3/8 inch (9,5mm) Sof-Lex Extra Thin (XT) Contouring and Polishing Discs – Super Fine
<b>Sof-Lex™ Mandrels</b>	
8695CA	2 – CA Mandrels
<b>Sof-Lex™ Finishing and Polishing Strips</b>	
1954	150 – 7 inches x 5/32 inch (178mm x 4mm) Sof-Lex Finishing and Polishing Strips – Coarse/Medium
1954N	100 - 7 inches x 5/64 inch (178mm x 2mm) Sof-Lex Finishing and Polishing Strips – Coarse/Medium, Narrow
1956	120 – 7 inches x 5/32 inch (178mm x 4mm) Sof-Lex Finishing and Polishing Strips – Fine/Super Fine

<b>Sof-Lex™ Contouring and Polishing Discs Kits</b>	
1980	<p>Sof-Lex Contouring and Polishing Disc Kit</p> <ul style="list-style-type: none"> <li>• 30 – 1/2 inch (12,7mm) Discs – Coarse</li> <li>• 30 – 1/2 inch (12,7mm) Discs – Medium</li> <li>• 30 – 1/2 inch (12,7mm) Discs – Fine</li> <li>• 30 – 1/2 inch (12,7mm) Discs – Super Fine</li> <li>• 30 – 3/8 inch (9,5mm) Discs – Coarse</li> <li>• 30 – 3/8 inch (9,5mm) Discs – Medium</li> <li>• 30 – 3/8 inch (9,5mm) Discs – Fine</li> <li>• 30 – 3/8 inch (9,5mm) Discs – Super Fine</li> <li>• 1 – Mandrel</li> </ul>
2380	<p>Sof-Lex Extra Thin (XT) Contouring and Polishing Disc Kit</p> <ul style="list-style-type: none"> <li>• 30 – 1/2 inch (12,7mm) XT Discs – Coarse</li> <li>• 30 – 1/2 inch (12,7mm) XT Discs – Medium</li> <li>• 30 – 1/2 inch (12,7mm) XT Discs – Fine</li> <li>• 30 – 1/2 inch (12,7mm) XT Discs – Super Fine</li> <li>• 30 – 3/8 inch (9,5mm) XT Discs – Coarse</li> <li>• 30 – 3/8 inch (9,5mm) XT Discs – Medium</li> <li>• 30 – 3/8 inch (9,5mm) XT Discs – Fine</li> <li>• 30 – 3/8 inch (9,5mm) XT Discs – Super Fine</li> <li>• 1 – Mandrel</li> </ul>
2385P	<p>Sof-Lex Finishing and Polishing Systems Kit</p> <ul style="list-style-type: none"> <li>• 30 – 1/2 inch (12,7mm) Discs – Coarse</li> <li>• 30 – 1/2 inch (12,7mm) Discs – Medium</li> <li>• 30 – 1/2 inch (12,7mm) Discs – Fine</li> <li>• 30 – 1/2 inch (12,7mm) Discs – Super Fine</li> <li>• 30 – 3/8 inch (9,5mm) Discs – Coarse</li> <li>• 30 – 3/8 inch (9,5mm) Discs – Medium</li> <li>• 30 – 3/8 inch (9,5mm) Discs – Fine</li> <li>• 30 – 3/8 inch (9,5mm) Discs – Super Fine</li> <li>• 30 – 1/2 inch (12,7mm) XT Discs – Coarse</li> <li>• 30 – 1/2 inch (12,7mm) XT Discs – Medium</li> <li>• 30 – 1/2 inch (12,7mm) XT Discs – Fine</li> <li>• 30 – 1/2 inch (12,7mm) XT Discs – Super Fine</li> <li>• 30 – 3/8 inch (9,5mm) XT Discs – Coarse</li> <li>• 30 – 3/8 inch (9,5mm) XT Discs – Medium</li> <li>• 30 – 3/8 inch (9,5mm) XT Discs – Fine</li> <li>• 30 – 3/8 inch (9,5mm) XT Discs – Super Fine</li> <li>• 15 – 7 inches x 5/32 inch (178mm x 4mm) Strips – Coarse/Medium</li> <li>• 15 – 7 inches x 5/64 inch (178mm x 2mm) Narrow Strips – Coarse/Medium</li> <li>• 15 – 7 inches x 5/32 inch (178mm x 4mm) Strips – Fine/Super Fine</li> <li>• 1 – Mandrel</li> </ul>



## CE ATITIKTIES DEKLARACIJA

Medicinos prietaisų 93/42/EEC Direktyvos VII priedas

**Gamintojas:** 3M Center, Bldg 260-2A-11, St.Paul, MN 55144 USA

**Gamintojo adresas:** 3M ESPE Dental Products  
2111 McGaw Avenue, Irvine, California 92614, USA

**ISO Sertifikato Nr.:** G1 11 11 45735 020  
**Galioja iki:** 2016-12-03  
**GMDN kodas:** 35807 – Diskai, odontologiniai abrazyvai  
35702 – Abrazyvinės juostelės, odontologinės  
35170 – Mandrelos, abrazyvinės  
**UMDNS kodas:** 16-201 – Poliravimo sistema  
11-282 – Šlifavimo ratukas

**Produkto grupė:** Abrazyvai  
**Pavadinimas:** **Produktai:** - Sof-Lex™ Šlifavimo ir poliravimo diskeliai  
- Sof-Lex™ Galutinio užbaigimo ir poliravimo diskeliai  
- Sof-Lex™ Galutinio užbaigimo ir poliravimo juostelės  
- Sof-Lex™ Labai ploni šlifavimo ir poliravimo diskeliai  
- Sof-Lex™ Galutinio užbaigimo ir poliravimo sistemos  
- Sof-Lex™ Mandrelos  
**CE ženklo data:** 1994 m. rugpjūčio mėn.  
**EC Produkto klasė:** I pagal Medicinos prietaisų IX priedą

### Atitikties deklaracija

3M ESPE Dental Products skelbia, kad priemonės išvardintos papildomoje priemonių lentelėje atitinka 1993m. liepos mėn., redaguotos 2007/47/EC, medicinos prietaisų 93/42/EEC EC Direktyvos būtinus reikalavimus, bei atitinka ISO 13485:2003, Medicinos prietaisų – kokybės valdymo sistemų – kontrolės reikalavimus, pagal Europos sąjungos medicinos prietaisų reikalavimus, patvirtintus Europos sąjungos Medicinos prietaisų reikalavimus.

3M ESPE Dental Products paskyrė 3M Deutschland GmbH kaip patvirtintą atstovą ES šalyse.

3M Deutschland GmbH  
Carl-Schurz-Strasse 1, 41453 Neuss, Germany (Vokietija)

2012-01-20

**Produkto grupė:** Abrazyvai  
**Produkto aprašymas:** Žiūrėti apačioje  
**EC produkto klasė (pagal IX priedo taisykles):** I klasė Medicinos prietaisų IX priedo 5 punktas  
**Pavadinimas:** - Sof-Lex™ Šlifavimo ir poliravimo diskeliai  
- Sof-Lex™ Galutinio užbaigimo ir poliravimo diskeliai  
- Sof-Lex™ Galutinio užbaigimo ir poliravimo juostelės  
- Sof-Lex™ Labai ploni šlifavimo ir poliravimo diskeliai  
- Sof-Lex™ Galutinio užbaigimo ir poliravimo sistemos  
- Sof-Lex™ Mandrelos

Katalogo/ užsakymo numeris	Aprašymas	
	<b>Sof-Lex™ Šlifavimo ir poliravimo diskeliai</b>	
8691C	12,7 mm Sof-Lex Šlifavimo ir poliravimo diskeliai - Šiurkštūs	
8691F	12,7 mm Sof-Lex Šlifavimo ir poliravimo diskeliai - Vidutiniai	
8691M	12,7 mm Sof-Lex Šlifavimo ir poliravimo diskeliai - Švelnūs	
8691SF	12,7 mm Sof-Lex Šlifavimo ir poliravimo diskeliai – Labai švelnūs	
8690C	9,5 mm Sof-Lex Šlifavimo ir poliravimo diskeliai - Šiurkštūs	
8690M	9,5 mm Sof-Lex Šlifavimo ir poliravimo diskeliai - Vidutiniai	
8690F	9,5 mm Sof-Lex Šlifavimo ir poliravimo diskeliai - Švelnūs	
8690SF	9,5 mm Sof-Lex Šlifavimo ir poliravimo diskeliai – Labai švelnūs	
	<b>Sof-Lex™ Labai ploni šlifavimo ir poliravimo diskeliai</b>	
8692C	12,7 mm Labai ploni (XT) šlifavimo ir poliravimo diskeliai - Šiurkštūs	
8692M	12,7 mm Labai ploni (XT) šlifavimo ir poliravimo diskeliai - Vidutiniai	
8692F	12,7 mm Labai ploni (XT) šlifavimo ir poliravimo diskeliai - Švelnūs	
8692SF	12,7 mm Labai ploni (XT) šlifavimo ir poliravimo diskeliai – Labai švelnūs	
8693C	9,5 mm Labai ploni (XT) šlifavimo ir poliravimo diskeliai - Šiurkštūs	
8693M	9,5 mm Labai ploni (XT) šlifavimo ir poliravimo diskeliai - Vidutiniai	
8693F	9,5 mm Labai ploni (XT) šlifavimo ir poliravimo diskeliai - Švelnūs	
8693SF	9,5 mm Labai ploni (XT) šlifavimo ir poliravimo diskeliai – Labai švelnūs	
	<b>Sof-Lex™ Mandrelos</b>	
8695CA	2 – CA Mandrelos	
	<b>Sof-Lex™ Galutinio užbaigimo ir poliravimo juostelės</b>	
1954	178 mm x 4 mm Galutinio užbaigimo ir poliravimo juostelės – Šiurkščios/Vidutinės	
1954N	178 mm x 2 mm Galutinio užbaigimo ir poliravimo juostelės – Šiurkščios/Vidutinės, Siauros	
1956	178 mm x 4 mm Galutinio užbaigimo ir poliravimo juostelės – Švelnios/Labai švelnios	
	<b>Sof-Lex™ Šlifavimo ir poliravimo diskelių rinkiniai</b>	
1980	Sof-Lex Šlifavimo ir poliravimo diskelių rinkinys - 12,7 mm diskeliai – šiurkštūs - 12,7 mm diskeliai – vidutiniai - 12,7 mm diskeliai – švelnūs - 12,7 mm diskeliai – labai švelnūs	- 9,5 mm diskeliai – šiurkštūs - 9,5 mm diskeliai – vidutiniai - 9,5 mm diskeliai – švelnūs - 9,5 mm diskeliai – labai švelnūs - 1 mandrela
2380	Sof-Lex Labai plonų šlifavimo ir poliravimo diskelių rinkinys - 12,7 mm XT diskeliai – šiurkštūs - 12,7 mm XT diskeliai – vidutiniai - 12,7 mm XT diskeliai – švelnūs - 12,7 mm XT diskeliai – labai švelnūs	- 9,5 mm XT diskeliai – šiurkštūs - 9,5 mm XT diskeliai – vidutiniai - 9,5 mm XT diskeliai – švelnūs - 9,5 mm XT diskeliai – labai švelnūs - 1 mandrela
2385P	Sof-Lex Galutinio užbaigimo ir poliravimo sistemų rinkinys - 12,7 mm diskeliai – šiurkštūs - 12,7 mm diskeliai – vidutiniai - 12,7 mm diskeliai – švelnūs - 12,7 mm diskeliai – labai švelnūs	- 9,5 mm diskeliai – šiurkštūs - 9,5 mm diskeliai – vidutiniai - 9,5 mm diskeliai – švelnūs - 9,5 mm diskeliai – labai švelnūs
	- 12,7 mm XT diskeliai – šiurkštūs - 12,7 mm XT diskeliai – vidutiniai - 12,7 mm XT diskeliai – švelnūs - 12,7 mm XT diskeliai – labai švelnūs	- 9,5 mm XT diskeliai – šiurkštūs - 9,5 mm XT diskeliai – vidutiniai - 9,5 mm XT diskeliai – švelnūs - 9,5 mm XT diskeliai – labai švelnūs
	- 178 mm x 4 mm Galutinio užbaigimo ir poliravimo juostelės – Šiurkščios/Vidutinės - 178 mm x 2 mm Galutinio užbaigimo ir poliravimo juostelės – Šiurkščios/Vidutinės, Siauros - 178 mm x 4 mm Galutinio užbaigimo ir poliravimo juostelės – Švelnios/Labai švelnios - 1 mandrela	

  
 VERTIMAS TEISINGAS  
  
 UAB „Plandent“ direktorius Andrius Preišegalavičius

## EG-KONFORMITÄTSERKLÄRUNG

entsprechend Anhang II der Richtlinie 93/42/EWG des Rates vom 14. Juni 1993 über  
Medizinprodukte

## EC ATTESTATION OF CONFORMITY

according to annex II of the Council Directive 93/42/EEC of June 14<sup>th</sup>, 1993 concerning  
medical devices

Wir: <b>Coltene/Whaledent GmbH + Co. KG</b> Raiffeisenstr. 30 D-89129 Langenau	We: <b>Coltene/Whaledent GmbH + Co. KG</b> Raiffeisenstr. 30 D-89129 Langenau
erklären in alleiniger Verantwortung, daß die Produkte:  Bezeichnung: Wurzelkanalfüllungsmaterialien Wurzelkanalbürsten Papierspitzen Absaugkanülen Sterile Mulltupfer Chirurgisches Nahtmaterial (nicht resorbierbar) Arzneimittelhaltige Retraktionsfäden Arzneimittelhaltige Gelatineschwämmchen Verbandmittel mit Jodoform Tamponadebinde Wurzelkanalfeilen Dentale rotierende Handstücke mit Bohrantrieb	declare under our sole responsibility that the products:  Name:  Root canal filling materials Root canal brushes Paper points Aspirator tips Sterile gauze swabs Surgical sutures (non-resorbable) Medicated retraction cords Medicated gelatin sponges Bandage with Iodoform Tamponade bandage Root canal files Dental rotary handpieces with engine
den einschlägigen Bestimmungen des Anhangs II der Richtlinie 93/42/EWG des Rates vom 14. Juni 1993 über Medizinprodukte entspricht.	meets the provisions of annex II of the Council Directive 93/42/EEC of June 14 <sup>th</sup> , 1993 concerning medical devices which apply to it.
Benannte Stelle: medical device certification GmbH Kriegerstraße 6 70191 Stuttgart Germany Nummer der Benannten Stelle: 0483	Notified Body: medical device certification GmbH Kriegerstraße 6 70191 Stuttgart Germany Number of the Notified Body: 0483
Diese EG-Konformitätsbescheinigung ist gültig bis 30.07.2018.	This EC-Attestation of conformity is valid until 30.07.2018.

Langenau, 18.07.13



**Barbara Wellbrock**  
Director QM/QC/RA

Coltene/Whaledent GmbH + Co. KG  
P.O.Box 1150, Raiffeisenstraße 30, D-89129  
Langenau/Gemany  
Tel.: ++49-7345-805-0  
<http://www.coltenewhaledent.de>

**coltene**   
**whaledent**®

## **EC ATITIKTIES DEKLARACIJA**

*Pagal 1993 07 14 medicinos prietaisų 93/42/EEC Tarybos Direktyvą annex II  
įtr. 2007/47/EU*

Mes:

*Coltene/Whaledent GmbH + Co. KG  
Raiffeisenstr. 30, D-89129 Langenau/Gemany*

**Būdami atsakingi skelbiame, kad produktai:**

Šaknų kanalų užpildymo medžiagos  
Šaknų kanalų šepetėliai  
Popieriniai kaiščiai  
Atsiurbėjai  
Sterilūs marlės tamponai  
Chirurginiai siūlai (netirpstantys)  
Medicininiai retrakciniai siūlai  
Medicininės želatinos kempinė  
Tvarsčiai su jodoformu  
Tvarsčiai  
Šaknų kanalų failai  
Odontologiniai sukamieji antgaliai su varikliu

*Atitinka 1993 07 14 medicinos prietaisų 93/42/EEC Tarybos Direktyvos annex II įtr.  
2007/47/EU reikalavimus*

*Notifikavimo įstaiga:        medicinos prietaisų sertifikavimas GmbH  
Kriegerstraße 6, 70191 Stuttgart, Vokietija*

*Notifikavimo numeris:    0483*

*Ši EC-Atitikties deklaracija galioja iki 2018-07-30.*

*Langenau 2013-07-18  
Barbara Wellbrock*

  
VERTIMAS TEISINGAS

UAB „Plandent“ direktorius Andrius Preišegalavičius

**EG-KONFORMITÄTSBESCHEINIGUNG**entsprechend Anhang VII der Richtlinie 93/42/EWG des Rates vom 14. Juni 1993 über  
Medizinprodukte**EC ATTESTATION OF CONFORMITY**according to annex VII of the Council Directive 93/42/EEC of June 14<sup>th</sup>, 1993 concerning  
medical devices

<b>Wir:</b> <b>Coltène/Whaledent GmbH + Co. KG</b> Raiffeisenstr. 30 D-89129 Langenau Germany	<b>We:</b> <b>Coltène/Whaledent GmbH + Co. KG</b> Raiffeisenstr. 30 D-89129 Langenau Germany
---	--

erklären in alleiniger Verantwortung, daß die Produkte: Bezeichnung: Abdecktücher Artikulationspapier Artikulationsseide Endodontikinstrumente Endodontikinstrument-Organisierer Handschuhe Kältespray Klemmpinzetten Kofferdam Kofferdam Klammern Kofferdam Klammerzange Kofferdam Lochzange Kofferdam Rahmen Kompressionshütchen Kopfschutztaschen Marker Mulltupfer Mundschutz Occlusionsfolie und Zubehör Patientenumhänge Polierstreifen Retraktionsfäden und Zubehör Schaumstoffpellets Schlauchüberzüge Schleifstreifen Shimstock-Folie Stahlmatrizenband Stahlseparierstreifen Wattekügelchen Watterollen Wurzelkanal-Tips	declare under our sole responsibility that the products: Name: Covers Articulating paper Articulating silk Endodontic instruments Endodontic organizer Gloves Cold spray Clamping tweezers Dental dam Dental dam clamps Dental dam clamp forceps Dental dam punch Dental dam frames Compression caps Headrest covers Markers Gauze swabs Face masks Occlusion foils and accessories Patient bibs Polishing strips Retraction cords and accessories Foam plastic pellets Sleeves for tubes Abrasive strips Shimstock-foil Steel matrix bands Steel separating strips Cotton pellets Cotton rolls Root canal tips
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den einschlägigen Bestimmungen des Anhangs VII der Richtlinie 93/42/EWG des Rates vom 14. Juni 1993 über Medizinprodukte entspricht.	<i>meets the provisions of annex VII of the Council Directive 93/42/EEC of June 14<sup>th</sup>, 1993 concerning medical devices which apply to it.</i>
Diese EG-Konformitätsbescheinigung ist gültig bis 30.07.2018.	<i>This EC-Attestation of conformity is valid until 30.07.2018.</i>

Langenau, 22.11.13

  
.....  
**Barbara Wellbrock**  
Director QM/QC/RA

Coltene/Whaledent GmbH + Co. KG  
P.O.Box 1150, Raiffeisenstraße 30, D-89129 Langenau/Gemany  
Tel.: ++49-7345-805-0  
<http://www.coltenewhaledent.de>

## **EC ATITIKTIES DEKLARACIJA**

*Pagal 1993 07 14 medicinos prietaisų 93/42/EEC Tarybos Direktyvą annex VII įtr.  
2007/47/EU*

Mes:

*Coltene/Whaledent GmbH + Co. KG*  
Raiffeisenstr. 30, D-89129 Langenau/Gemany

Būdami atsakingi skelbiame, kad produktai:

Uždangalai  
Atsiurbėjų adapteriai  
Artikuliacinis popierius  
Artikuliacinis šilkas  
Endodontiniai instrumentai  
Pirštinės  
Šaldymo aerozoliai  
Uždėjimo replės  
Dantų koferdamo guma  
Dantų koferdamo žiedai  
Dantų koferdamo gumos pradūrėjas  
Dantų koferdamo rėmeliai  
Užspaudimo kepurėlės  
Galvos atlošo uždangalai  
Markeriai  
Seilių ritinėliai  
Veido apsauginės kaukės  
Sąkandžio folijos ir priedai  
Pacientų apsauginės servetėlės  
Poliravimo juostelės  
Retrakciniai siūlai ir priedai  
Putplasčio ritinėliai  
Apsauginiai apvalkalai vamzdeliams  
Abrazyvinės juostelės  
Shimstock folija (sąkandžiui)  
Plieninės matricos  
Plieninės atskyrimo juostelės  
Lignino ritinėliai

*Atitinka 1993 07 14 medicinos prietaisų 93/42/EEC Tarybos Direktyvos annex VII įtr.  
2007/47/EU reikalavimus*

*Ši EC-Atitikties deklaracija galioja iki 2018-07-30.*

*Langenau 2013-11-22*  
*Barbara Wellbrock*

VERTIMAS TEISINGAS

UAB „Plandent“ direktorius Andrius Preišgalavičius





## DECLARATION OF CONFORMITY

We

**Plandent Oy,  
Asentajankatu 6,  
FIN-00880 Helsinki  
Finland**

ensures and declares that the products

Orbis Perforated Diamond Strips

are Class I products according to Annex IX of Directive 93/42/EEC and are manufactured in conformity with the provisions of the Council Directive 93/42/EEC and the technical documentation referred in Annex VII, Section 3 and Annex V as amended by 2007/47/EC thus, meeting requirements of the Medical Device Directive.

Helsinki, 2015-8-9

  
Senja Tynkkynen  
Director Sourcing



## ATITIKTIES DEKLARACIJA

Mes

**Plandent Oy,  
Asentajankatu 6,  
FIN-00880  
Helsinki Finland  
(Suomija)**

užtikriname ir pareiškiame, kad produktas

Orbis perforuotos deimantinės juostelės

pagal 93/42/EEB direktyvą yra I klasės produktai ir yra pagamintos laikantis Tarybos direktyvos 93/42/EEB sąlygų ir techninė dokumentacija, nurodyta VII priedo 3 skirsnyje ir V priedo pakeitimuose, atliktuose 2007/47/EB atitinka Medicinos prietaisų direktyvos reikalavimus.

Helsinkis, 2015-08-09

Senja Tynkkynen  
Tiekimo direktorė

VERTIMAS TEISINGAS

UAB „Plandent“ direktorius Andrius Preišgalavičius



## EG-Zertifikat / EC-Certificate

**gem. 93/42/EWG Anhang V / acc. 93/42/EEC Annex V**

Hiermit wird bescheinigt, dass die Firma / *This certifies, that the company*

**ORBIS Dental Handelsgesellschaft mbH**  
**Schuckertstraße 21**  
**48153 Münster**  
**Deutschland**

für die Produkte / die Kategorie: Liste der Produkte siehe Anlage 1  
*for the products / product category: List of products see annex 1*

***Dentalprodukte, sterile Mulltupfer***  
***Dental products, sterile gauze swab***

ein Qualitätssicherungssystem für die Herstellung der genannten Produkte nach Maßgabe des Anhang V der Richtlinie 93/42/EWG anwendet. Die Übereinstimmung mit den Anforderungen wurde in einem Audit nachgewiesen. Zusätzlich zur CE-Kennzeichnung muss die Kennnummer der Benannten Stelle angebracht werden. Die Gültigkeit dieses Zertifikats beruht auf der Aufrechterhaltung des Qualitätssicherungssystems in Übereinstimmung mit den Anforderungen der Richtlinie und seiner Überwachung durch die Benannte Stelle gem. Anhang V Abschnitt 4. Das Zertifikat ist unter keinen Umständen übertragbar.

*has established a quality system for production acc. to the requirements of Annex V of the directive 93/42/EEC. Conformity with the requirements was proved within an audit. Additional to the CE-marking the notification number of the Notified Body has to be affixed. The validity of this certificate is based on the maintenance of the quality system in accordance with the requirements of the directive and its surveillance by the Notified Body according Annex V section 4. The certificate may not be transferred under any circumstances.*

Reg.-Nr. / Reg.-No. 44 235 121859  
Bericht Nr. / Report No. 3510 8294

Gültigkeit / Validity  
von / from 2013-01-09  
bis / until 2017-09-30  
Edition 1



Zertifizierungsstelle für Medizinprodukte  
*Certification body for medical devices*

Essen, 2013-01-09

TÜV NORD CERT GmbH    Langemarckstraße 20    45141 Essen    www.tuev-nord-cert.de    medical@tuev-nord.de

Benannte Stelle Kenn-Nr. 0044 / *Notified Body ID. No. 0044*

Benannt durch / *Notified by*  
Anerkannt durch / *Recognized by*  
ZLG, ZLS: [www.zlg.de](http://www.zlg.de)

# ANLAGE / ANNEX

Anlage 1, Blatt 1 von 1  
Annex 1, page 1 of 1

**Reg.-Nr. / Reg. No. 44 235 121859**

Produkte der Klasse Is  
*Products of class Is*

## **Sterile Mulltupfer**

**Anmerkung:** Für Produkte der Klasse I steril beschränkt sich das Zertifizierungsverfahren auf die Aspekte der Herstellung unter Sicherstellung und Überwachung der sterilen Bedingungen.

**Note:** *For products of class I sterile the certification process is restricted to the aspects of manufacture concerned with securing and maintaining sterile conditions.*

Produkte der Klasse IIa  
*Products of class IIa*

- **Produkte zur Desinfektion von Medizinprodukten**
- **zahnärztliche Werkstoffe**
- **nichtaktive zahnärztliche Instrumente**

Bericht Nr. / Report No. 3510 8294

Gültigkeit / Validity  
von / from 2013-01-09  
Edition 1

A handwritten signature in blue ink, appearing to be a stylized 'J' followed by a horizontal line.

Zertifizierungsstelle für Medizinprodukte  
*Certification body for medical devices*

Essen, 2013-01-09

TÜV NORD CERT GmbH    Langemarckstraße 20    45141 Essen    [www.tuev-nord-cert.de](http://www.tuev-nord-cert.de)    [medical@tuev-nord.de](mailto:medical@tuev-nord.de)

Benannte Stelle Kenn-Nr. 0044 / *Notified Body ID. No. 0044*

**Benannt durch / Notified by**  
**Anerkannt durch / Recognized by**  
ZLG, ZLS: [www.zlg.de](http://www.zlg.de)

## EC – Sertifikatas

### Pagal 93/42/EEC V priedą

Tai patvirtina, kad kompanija

**ORBIS Dental Handelsgesellschaft mbH**

**Schuckertsraße 21**

**48153 Münster**

**Vokietija**

Produktams / produktų kategorijoms

(Produktų sąrašą žiūrėti 1 priede)

### **Odontologiniai produktai, sterilūs marlės tamponai**

*Įdiegė kokybės sistemą produkcijai pagal 93/42/EEC direktyvos V priedą. Atitiktis su reikalavimais buvo įrodyta audito. Papildomai prie CE žymėjimo Notifikavimo įstaiga suteikė notifikacijos numerį. Šio sertifikato galiojimas remiasi kokybės sistemos palaikymu pagal direktyvos reikalavimus ir yra Notifikavimo įstaigos priežiūroje pagal V priedo 4 dalį. Dėl tam tikrų sąlygų Sertifikatas gali būti neperduodamas.*

Reg.Nr. 44 235 121859

Protokolo Nr. 3510 8294

Galioja nuo 2013-01-09

Galioja iki 2017-09-30

Leidimas 1

*Medicinos prietaisų sertifikavimo įstaiga*

Essen, 2013-01-09

TÜV NORD CERT GmbH, Langemarckstr. 20, D-45141 Essen

Notifikavimo įstaigos ID Nr. 0044

1 priedas, 1 puslapis (viso 1 puslapis)

**Reg.Nr. 44 235 121859**

*Produktų klasė I*

### **Sterilūs marlės tamponai**

Pastaba: I klasės produktams sterilumo sertifikavimo procesas apsiriboja tik toms gamybos grandims, kurių metu užtikrinamas sterilumas.

### **Ia klasės produktai**

- **Produktai, skirti medicinos prietaisų dezinfekcijai**
- **odontologinės medžiagos**
- **neaktyvūs odontologijos instrumentai**

Protokolo Nr. 3510 8294

Galioja nuo 2013-01-09

Leidimas 1

*Medicinos prietaisų sertifikavimo įstaiga*

Essen, 2013-01-09

TÜV NORD CERT GmbH, Langemarckstr. 20, D-45141 Essen

Notifikavimo įstaigos ID Nr. 0044

VERTIMAS TEISINGAS

UAB „Plandent“ direktorius Andrius Preišegalavičius





## DECLARATION OF CONFORMITY

Ultradent Products, Inc. has evaluated the following product by using the Conformity Assessment Procedure of Annex VII of the Medical Device Directive 93/42/EEC, as amended by 2007/47/EEC:

### Syringe, Compule, Luer Lock Cap, Syringe Covers and Barrier Sleeve

and confirms that the product is compliant with the Essential Requirements of Annex I of the Medical Device Directive 93/42/EEC. Technical documentation is located in the Regulatory Affairs Department.

This product system is classified as Class I medical device according to the Medical Device Directive 93/42/EEC, Section III Classification, 2.1 Rule 5.

**UMDNS Code:** 13938, Dental syringes  
11160, Dental hand instrument  
**GMDN Code:** 45051, Applicator dental  
16825, Syringe tip cap  
17977, Cover handgrip

*Matt Buck*  
Matt Buck  
Regulatory Affairs Specialist

12/01/10  
Date

State of Utah, County of Salt Lake  
Subscribed and sworn to before me  
this 1 day of December 2010

*Diane E. Rogers*  
Diane Rogers, Notary Public



This document is in force as long as the following ISO certificates are valid:

EC Certificate 44 232 06 342325-001  
EN ISO 13485 :2007 44 221 06 342325-001

TF044.2



## ATITIKTIES DEKLARACIJA

Utradent Products, Inc. įvertino šiuos produktus naudodamiesi Atitikties įvertinimo procedūra pagal Medicinos prietaisų direktyvą 93/42/EEB VII priedą ir 2007/47/EEB pakeitimus:

### Švirkštai, sujungimai, Luer Lock cap dangteliai, švirkštų uždengimai ir apsaugos

ir patvirtina, kad šie produktas atitinka medicinos prietaisų direktyvos 93/42/EEB I priedo pagrindinius reikalavimus. Techninė dokumentacija yra laikoma Reguliavimo reikalų departamente.

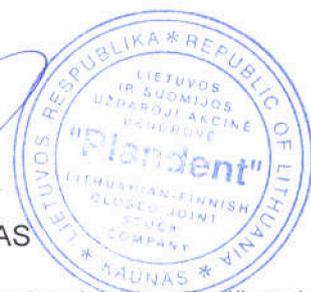
Ši produkto sistema yra klasifikuojama kaip I klasės medicinos prietaisas pagal Medicinos prietaisų direktyvos 93/42/EEB III dalies klasifikacija 2.1, 5 taisyklė.

**UMDNS kodas:** 13938, Odontologiniai švirkštai  
11160, Odontologiniai rankiniai instrumentai  
**GMDN kodas:** 45051, Odontologinis aplikatorius  
16825, Švirkštų antagliukai  
17977, Rankenos uždengimas

Matt Buck  
Reguliavimo reikalų specialistas

2010/01/12  
Data

Šis dokumentas galioja kol galio ISO sertifikatai:  
EB Sertifikatas 44 232 06 342325-001  
EN ISO 13485:2007 44 221 06 342325-001

  
VERTIMAS TEISINGAS  
  
UAB „Plandent“ direktorius Andrius Preišgalavičius



# DECLARATION OF CONFORMITY

Ultradent Products, Inc. has evaluated the following product by using the Conformity Assessment Procedure of Annex VII of the Medical Device Directive 93/42/EEC, as amended by 2007/47/EEC:

## Tips

and confirms that the product is compliant with the Essential Requirements of Annex I of the Medical Device Directive 93/42/EEC. Technical documentation is located in the Regulatory Affairs Department.

This product system is classified as Class I medical device according to the Medical Device Directive 93/42/EEC, Section III Classification 2.1, Rule 5.

**UMDNS Code:** 11160, Dental hand instrument

**GMDN Code:** 35917, Suction tip, general purpose, single-use  
16825, Syringe tip cap



Matt Buck  
Regulatory Affairs Specialist



Date

State of Utah, County of Salt Lake  
Subscribed and sworn to before me  
this 28 day of June 2012

  
Diane Rogers, Notary Public



This document is in force as long as the following ISO certificates are valid:

EC Certificate 44 232 06 342325-001  
EN ISO 13485 44 221 06 342325-001



## ATITIKTIES DEKLARACIJA

Ultradent Products, Inc. įvertino šiuos produktus naudodamiesi Atitikties įvertinimo procedūra pagal Medicinos prietaisų direktyvą 93/42/EEB VII priedą ir 2007/47/EEB pakeitimus:

### Antgaliukai (Tips)

ir patvirtina, kad šie produktas atitinka medicinos prietaisų direktyvos 93/42/EEB I priedo pagrindinius reikalavimus. Techninė dokumentacija yra laikoma Reguliavimo reikalų departamente.

Ši produkto sistema yra klasifikuojama kaip I klasės medicinos prietaisas pagal Medicinos prietaisų direktyvos 93/42/EEB III dalies klasifikacija 2.1, 5 taisyklė.

**UMDNS kodas:** 11160, Odontologiniai rankiniai instrumentai  
**GMDN kodas:** 35917, Atsiurbimo antgaliukai, bendrosios paskirties, vienkartinio naudojimo  
16825, Švirkštų antgaliukai

Matt Buck  
Reguliavimo reikalų specialistas

2012/06/28  
Data

Šis dokumentas galioja kol galio ISO sertifikatai:  
EB Sertifikatas 44 232 06 342325-001  
EN ISO 13485 44 221 06 342325-001

VERTIMAS TEISINGAS

UAB „Plandent“ direktorius Andrius Preišegalavičius

## Declaration of Conformity

**Manufacturer:** Panadent Corporation

**Address:** 580 South Rancho Ave  
Colton, CA 92324  
USA

The medical device:

Product Code	Product Name
4405	Bite Tab Compound Disks

Classification: Class I Short Term per Annex IX Definition 1.1 and Rule 5 of the 93/42/EEC directive as amended by 2007/47/EC.

Conformity assessment procedure according to MDD directive - Annex VII per Article 11-5

We, Panadent Corporation, declare and ensure with sole responsibility, that the above mentioned products meet the provisions of the Council Directive 93/42/EEC, Medical Device Directive as amended by 2007/47/EC which applies to them.

**Harmonized Standards:** ISO 13485:2003  
ISO 14971:2012  
EN 980:2008  
EN 1041:2008

**Additional information:**

**Notified Body:** Nemko AS

**EC Representative:** MDSS GmbH  
**EC Representative Address:** Schiffgraben 41  
30175 Hannover  
Germany

Signature of the Authorized Representative:

  
\_\_\_\_\_  
Panadent Corporation

09/11/2014  
\_\_\_\_\_  
Date

**ATITIKTIES DEKLARACIJA****Gamintojas: Panadent Corporation**Adresas: 580 South Rancho Ave  
Colton, CA 92324  
USA (JAV)

Medicinos prietaisas:

<b>Produkto kodas</b>	<b>Produkto pavadinimas</b>
4405	Bite Tab Compound Disks

Klasifikacija: I klasė pagal 93/42/EEB direktyvos IX priedo 1.1 apibrėžimą ir 2007/47/EEB pakeitimus.

Atitikties įvertinimo procedūra pagal MPD direktyvos VII priedo 11-5 straipsnį

Mes, Panadent Corporation, būdami atsakingi skelbiame ir tvirtiname, kad viršuje minimi produktai atitinka Medicinos prietaisų 93/42/EEB tarybos direktyvos nuostatas bei joms taikomus 2007/47/EEB pakeitimus.

Suderinti standartai: ISO 13485:2003  
ISO 14971:2012  
EN 980:2008  
EN 1041:2008

Papildoma informacija:

Notifikavimo įmonė: Nemko AS

EB atstovas: MDSS GmbH  
EB atstovo adresas: Schiffgraben 41  
30175 Hannover  
Germany (Vokietija)

Parašas autorizuoto atstovo:

2014-11-09

VERTIMAS TEISINGAS

UAB „Plandent“ direktorius Andrius Preišegalavičius





## Dental Laboratory and Surgery Materials

Issue: 1

Prepared by: J. A. Holder  
Date: 20<sup>th</sup> January 2015

### EC DECLARATION OF CONFORMITY

Associated Dental Products Limited hereby declare all devices on the attached Schedule of Devices conform with the appropriate requirements of the Medical Devices Directive 93/42/EEC as amended by EC Council Directive 2007/47/EC.

### DWS219 ALMINAX WAX 10 SHEETS 250G

Signed:.....*A Mook*.....

Date:.....*20/1/15*.....

QUALITY MANAGER



## Dental Laboratory and Surgery Materials

Leidimas: 1  
Paruošė: J.A.Holder  
Data: 2015 sausio 20d.

### EB ATITIKTIES DEKLARACIJA

Associated Dental Products Limited pareiškia, kad visi prietaisai pateiktame prietaisų išvardinime atitinka Medicinos prietaisų direktyvos 93/42/EEB bei 2007/47/EB tarybos direktyvos pakeitimų tinkamus reikalavimus.

### DWS219 ALMINAX WAX (vaškas) 10 SHEETS (lapelių) 250G

Parašas  
Kokybės vadovas

Data 2015 01 20

VERTIMAS TEISINGAS

UAB „Plandent“ direktorius Andrius Preišegalavičius



**Film/Sensor Holding Instruments/Disposables/Forceps<sup>1</sup>**  
**EC Declaration of Conformity**

**Reference:** See attached Schedule

**Manufacturer' Name:** DENTSPLY RINN  
**And Address:** 1212 Abbott Drive  
Elgin, Illinois 60123

**Authorized Representative:** DENTSPLY Limited  
**And Address:** Building 1, Aviator Park, Station Road  
Addlestone, Surrey KT15 2PG – UK

**Name/Description of product:** Film/Sensor Holding Instruments/Disposables  
**Device Description:** Dental X-ray film holder  
Dental Hand Instrument<sup>1</sup>

**GMDN code:** 16205, 31813<sup>1</sup>

**Classification:** Class I

**EC Declaration of Conformity**

The above listed medical devices conform to the requirements of Council Directive 93/42/EEC and meet the relevant Essential Requirements of Annex I. All devices are designed, manufactured, and tested and released for sale in accordance with the technical documentation.

This declaration is made on the basis of self certification by DENTSPLY RINN in accordance with Annex V of the Directive.

**Date of Issue** 07/24/1997

**Revision** H

**Signed:**

William T. Cousins  
William T. Cousins

**Rev. Date:**

May 31, 2011

**Title:**

Manager QA/RA

**DENTSPLY RINN Division maintains a quality management system that complies with the United States Food and Drug Administration 21 CFR Part 820, ISO 13485:2003 and the European Union Council Directive 93/42/EEC.**

## **SCHEDULE: Film/Sensor Holding Instruments/Disposables/Forceps**

Please refer to the attached declaration of conformity

### Film/Sensor Holding Instruments

XCP® and BAI Instruments

XCP-DS® Digital Sensor Holders

ENDORAY® II

UNI-BITE™

Uni-GripAR®

Snap-A-Ray®

Snap-A-Ray® Xtra

Snap-A-Ray® DS

Eezee-Grip® Universal Sensor Holder

Private Label Digital Sensor Holders

FLIP-RAY®

XCP-ORA®

VIPTM Film Holder

Cover Cozee®

EZ-Tab ® BW Tab for Sensors

XCP-DS® for Schick #1 V BW Elite

XCP-DS® for Schick #2 V BW Elite

XCP-ORA® Kit for SCHICK Elite

XCP-DS FIT™

### DISPOSABLES

XCP® and STABE® Bite Blocks

EZ-PROPTM MOUTH PROPS

Uni-Grip®

**Uni-Grip® 360 Universal Disposable Sensor Holder**

Rinn® Digital Sensor Loops

Rinn® Adhesive Bite-Wing Tabs

Rinn® Bite-Wing-Loops

Rinn® Digital Sensor Covers

E-Z Hold Articulating Paper Forceps<sup>1</sup>

## Filmučių (juostelių)/ Jutiklio laikymo instrumentų/Padėkliukų/ Žnyplių

### EB ATITIKTIES DEKLARACIJA

**Nuoroda**  
**Gamintojo pavadinimas**  
**Adresas:**

Žiūrėti pridėtame apraše  
DENTSPLY RINN  
1212 Abbott Drive  
Elgin, Illinois 60123

**Patvirtintas EB atstovo**  
**Pavadinimas ir adresas:**

DENTSPLY Limited  
Building 1, Aviator Park, Station Road  
Addlestone, Surrey KT15 2PG- UK

**Produkto pavadinimas/aprašymas:** Filmutės (juostelės)/ Jutiklio laikymo instrumentai/Padėkliukai  
**Prietaiso aprašymas:** Odontologiniai rentgeno juostelių laikikliai  
Odontologiniai rankiniai instrumentai

**GMDN kodas:** 16205, 31813

**Klasifikacija:** I klasė

#### EB Atitikties Deklaracija

Viršuje išvardinti medicinos prietaisai atitinka 93/42/EEB Tarybos direktyvos reikalavimus bei esminius I priedo reikalavimus. Visi prietaisai yra sukurti, pagaminti ir išbandyti bei išleisti pardavimui laikantis techninės dokumentacijos.

Ši deklaracija sukurta DENTSPLY rin sertifikavimo pagrindu pagal direktyvos V priedą.

**Išdavimo data** 1997/047/24

**Peržiūrėjimas** H

**Parašas:** William T.Cousins  
**Pareigos:** QA/RA vadovas

**Perž.data:** 2011-05-31

DENTSPLY RINN padalinys palaiko kokybės valdymo sistemą, kuri atitinka Jungtinių Valstijų maisto ir vaistų tvarkymo 21 CFR 820 dalį, ISO 13485:2003 ir Europos sąjungos Tarybos direktyvą 93/42/EEB.

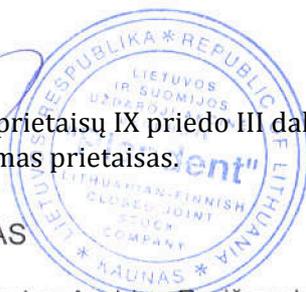
**Produkto klasifikacija**  
**Pagal IX priedą:**

Šis produktas klasifikuojamas pagal medicinos prietaisų IX priedo III dalies, 8 taisyklę kaip IIa klasės, ilgalaikis implantuojamas prietaisas.

1 puslapis iš 2

VERTIMAS TEISINGAS

UAB „Plandent“ direktorius Andrius Preišgalavičius



## APRAŠAS: **Filmutės (juostelės)/ Jutiklio laikymo instrumentai/Padėkliukai/ Žnyplės**

Prašome žiūrėti pridėtoje atitikties deklaracijoje

### **Filmutės (juostelės)/ Jutiklio laikymo instrumentai**

XCP® ir BAI instrumentai

XCP-DS® skaitmeniniai jutiklio laikikliai

ENDORAY® II

UNI-BITE™

Uni-Grip AR®

Snap-A-Ray®

Snap-A-Ray®Xtra

Snap-A-Ray®DS

Eezee-Grip®Universalus jutiklio laikiklis

Private Label Digital Sensor Holders (Privataus ženklo skaitmeniniai jutiklių laikikliai)

FLIP-RAY®

XCP-ORA®

VJPTM juostelių laikiklis

Cover cozee®

EZ-Tab® BW Tab for Sensors

XCP-DS® for Schick #1 V BW Elite

XCP-DS®for Schick #2 V BW Elite

XCP-ORA® Rinkinys SCmCK Elite

XCP-DS FIT™

### **Padėkliukai**

XCP® ir STABE® Bite Blokeliai

EZ-PROP™ MOUTH PROPS

Uni-Grip®

**Uni-Grio® 360 Universalus vienkartinis Sensoriaus laikiklis**

Rinn® Digital Sensor Loops

Rinn® AdhesiveBite-Wing Tabs

Rinn® Bite-Wmg-Loops

Rinn® Digital Sensor Covers (skaitmeninis jutiklio apdangalas)

E-Z Hold Articulating Paper Forceps (artikuliacinio popieriaus žnyplės)

## EEC-CONFORMITY DECLARATION FOR MEDICAL PRODUCTS

### EC COUNCIL DIRECTIVE 93/42/EEC FOR MEDICAL PRODUCTS

**Manufacturer:** **LM-Instruments Oy**  
PL 88  
21601 Parainen  
Finland  
Tel. +358 2 4546 400  
Fax +358 2 4546 444

Herewith we declare that all the dental hand instruments we are producing as well as LM-ErgoHold3, LM-Cello, LM-Multiseptor, LM-CompoKnife and LM-Activator comply with the relevant regulations of the guideline 93/42/EEC for medical products. Our products are classified in class I.

Date, place: 27.08.2009, Parainen



Kari Lehtonen  
Development Manager



**MEDICINOS PRODUKTŲ EEC ATITIKTIES DEKLARACIJA**  
**ES TARYBOS MEDICINOS PRIEMONIŲ DIREKTYVA 93/42/EEC**

**Gamintojas:** **LM - Instruments Oy**  
PL 88  
21601 Parainen  
Suomija  
Tel.: +358 2 4546 400  
Faks.: + 358 2 4546 444

Šiuo raštu pareiškiame, kad visi mūsų gaminami rankiniai odontologiniai instrumentai, taip pat ir LM ErgoHold3, LM-Cello ir LM Multiseptor, LM CompoKnife ir LM Activator atitinka Medicinos priemonių Direktyvos 93/42/EEC reikalavimus. Mūsų produktai priskiriami I-ai klasei.

Data, vieta: 2009-08-27, Parainen

Kari Lehtonen  
Plėtros vadybininkas



## Document Detail

**Type:** 1450-RDT\_Dec of Conf  
**Document No.:** 1450-RDT\_Dec of Conf\_A35[6]  
**Title:** RDT-Declaration of conformity-A35  
**Comment:** A35-Declaration of conformity 05  
**Status:** CURRENT  
**Effective Date:** 26-Nov-2014

### Approvals

<u>Role</u>	<u>Sign-off By</u>	<u>Sign-off Date</u>
Maillefer-PMCS B4,D1-PPCS 803 Dir. AQ & AR REG administrator	Luc Etienne Glawdys Monnin	26-Nov-2014 8:28 am GMT 26-Nov-2014 8:33 am GMT

### Reference

<u>Document No.</u>	<u>Title</u>	<u>Relation</u>
<a href="#">1450-RDT_Summary_A35 [3]</a>	A35-Table of content	Master Document

CE declaration of conformity – Directive 93/42/EEC

**Class I products**

We

**MAILLEFER INSTRUMENTS HOLDING Sàrl**

(including the subsidiaries: Maillefer Instruments Trading sàrl, Maillefer Instruments Manufacturing sàrl, Maillefer Instruments Consulting sàrl)  
Trading as Dentsply Maillefer

Chemin du Verger 3, CH-1338 Ballaigues / Switzerland

declare on our own responsibility that the following products listed in enclosure F 711.208.E: **Reference List A35** (4 pages), version n°06.

Conform to the requirements of Council Directive 93/42/EEC and meet the Essential Requirements of Annex I. All devices are designed, manufactured, tested and released for sale in accordance with the technical documentation.

This declaration is based on the following data:

-  Technical documentation demonstrating the product conformity to the requirements of the directive and the edict.
-  Applied norms: Listed in **Reference list A35**
-  Conformity assessment procedure: Annex VII of the directive 93/42/EEC

Dentsply Maillefer's Division maintains a quality management system that complies with the United States Food and Drug Administration 21 CFR Part 820, EN ISO 13485:2012, the European Union Council Directive 93/42/EEC, and the Japanese PAL law.

Ballaigues, 24/11/2014

Management Representative



Luc Etienne

MAILLEFER INSTRUMENTS  
HOLDING SàrlEncl.

## EC Declaration of conformity - Reference list

Product or family designation: <b>Accessories</b>		
File number: <b>A35</b>		
Prepared by: <b>Yassin Rouhila</b>	Visa: <i>YRO</i>	Date: <i>24/11/2014</i>
Reviewed by: <b>Luc Etienne</b>	Visa: <i>[Signature]</i>	Date: <i>24.11.2014</i>

Document History:

Version	Date	Change summary
06	24.11.2014	Update of the reference's list.

Reference list:

Code	Réf.	Proprietary Name	Intended use	GMDN #	GMDN definition	Rule	Class	S/M	Standards
	A0197	Silicone-stops dispenser	A simple and fast way to put a silicone stop on a root canal instrument. When the white button is pressed, the stop appears and with the tip of the instrument it can easily be picked up.	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041.
	A0197A	Silicone stops	Accessory to adjust the working length.	38442	Unclassified	5-1 <sup>st</sup> hyphen	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.
R	A203M-1	Split-Kit M empty	Sterilizable kit which serves to store instruments.	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.
N	A203M-9	Split-Kit M empty	Sterilizable kit which serves to store instruments.	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.

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	A203V	Split-Kit empty	Sterilizable kit which serves to store instruments.	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.
D	A0207	Endo-Container Complete	Kit used to store Gutta-Percha-points, paper points or other instruments.	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.
D	A207T	Tube Endo-Container	Tube used to store Gutta-Percha-points, paper points or other instruments.	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.
	A219P	Plastic Mandrel	For manual use of the root canal instrumentation.	38442	Unclassified	5-1 <sup>st</sup> hyphen	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.
R	A0227	Metallic Split Part		38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.
R	A227T	Tube	Gutta-Percha-Points, Paper Points or other instrument storage.	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041.
	A259A	Cotton pellet	Device permitting addition of disinfectant solution.	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041.
	A265P	Endo-Stand	Sterilizable kit which serves to store instruments.	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.
	A0266	Endo Organizer Empty	Sterilizable kit for ProFile system.	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.
	A268P	Clean-Stand	Use to store instruments in a clean environment during endodontic treatment.	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.
	A268P-2	Clean-Stand Large Version	Use to store instruments in a clean environment during endodontic treatment.	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.

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D	A268S	Sponges Clean- Stand	Foam-rubber blocks for Clean- stand Ref. A268P.	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.
D	A268S-2	Sponges Clean- Stand Large Version	Foam-rubber blocks for Clean- stand Ref. A268P-2.	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.
	A272V	Sosio-box empty	Sterilizable kit which serves to store instruments.	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.
	A274V	Flexokit empty	Sterilizable kit which serves to store instruments.	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.
	A0290	Inside Endo Prof. Bogopolsky	Plastic insert with holes for instruments and a foam block can be used in the Split kit.	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.
R	A0349	Multi-Stand	Sterilizable kit for Profile system.	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.
	A0362-9	ProFile Organizer Empty	Sterilizable kit for ProFile system.	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.
	A0413	ProTaper sequencer	Kit which serves to store instruments.	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.
R	A0482	Sticker GT obturator		38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041.
R	A0491	System GT organizer		38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.
R	A0582	Sticker GT series X Obturator		38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041.

## EC Declaration of conformity - Reference list

	A0591	GT series X organizer	Sterilizable kit which serves to store instruments.	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.
D	A1007-2	X-Smart AC Adapter	Use for power supply of endodontic motor.	38442	Unclassified	12	I	-	ISO, 13485, ISO 14971, EN 60601-1, EN 1640, ISO 11498, ISO 7494-1, ISO 15223-1.
D	A1007-4	X-Smart Foot Pedal	Use to control endodontic motor.	38442	Unclassified	12	I	-	ISO, 13485, ISO 14971, EN 60601-1, EN 1640, ISO 11498, ISO 7494-1, ISO 15223-1.
R	A1007-8	X-Smart Spray	Use for lubricating X-Smart.	15772	Lubricant, instrument	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041.
	A1409	NiTi manual handle adaptors	For manual use of the root canal instrumentation.	38442	Unclassified	5-1 <sup>st</sup> Hyphen	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041.
R	A1416	Sticker ProTaper Universal Obturator	Accessory to adapt Thermanprep oven to ProTaper Universal obturators.	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041.
	E0248	Sterilizable plastic box	Storage and sterilization of instruments.	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.
	E0249	Plastic insert for box E0248	Inserts for the storage of burs and Endo instruments in the sterilizable container Ref. E0248.	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.
	E0335	Box for FG	Kit for Burs	38442	Unclassified	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1.

**Dokumento detalės**

**Tipas:** 1450-RDT\_Atit Dekl  
**Dokumento Nr.:** 1450-RDT\_Atit Dekl\_A35[6]  
**Pavadinimas:** RDT-Atitikties deklaracija-A35  
**Komentarai** A35-Atitikties deklaracija 05  
**Statusas:** AKTUALUS  
**Įsigaliojimo data:** 2014-11-26

**Patvirtinimai****Vaidmuo**

Maillefer-PMCS B4,D1-PPCS 803 Dir. AQ & AR  
 REG administrator

**Pasirašyta**

Luc Etienne  
 Glawdys Monnin

**Pasirašymo data**

2014-11-26 8:28 GMT  
 2014-11-26 8:33 GMT

**Nuoroda****Dokumento Nr.**

1450-RDT\_Summary\_A35 [3]

**Pavadinimas**

A35-Table of content

**Ryšys**

Master Document



F 711.204

CE atitikties deklaracija – 93/42/EEB Direktyva  
 I Klasės produktai

Mes,

**MAILLEFER INSTRUMENTS HOLDING Sarl ir filialai**

(kartu su: Maillefer Instruments Trading Sarl, Maillefer Instruments Manufacturing Sarl,  
 Maillefer Instruments Consulting Sarl)  
 Nurodoma kaip Dentsply Maillefer

**Chemin du Verger 3, CH-1338 Ballaigues / Switzerland**

būdami atsakingi skelbiame, kad visi medicininiai prietaisai išvardinti priede F 711.208.E  
**Ref.sąrašas A35** (4 puslapiai), versija nr.06

Atitinka Tarybos direktyvos 93/42/EEB reikalavimus ir I priedo pagrindinius reikalavimus.  
 Visi prietaisai yra sukurti, pagaminti, išbandyti ir išleisti į rinką pagal techninę dokumentaciją.

Ši deklaracija remiasi sekančiais informacija:

- Techniniai dokumentai, įrodantys produktų atitikties direktyvos reikalavimus viršuje paminėtos direktyvos.
- Taikomos normos: I švardintos **Reference list A35**
- Atitikties įvertinimo procedūra: 93/42/EEB direktyvos VII priedas
- 

Dentsply Maillefer padalinys palaiko kokybės valdymo sistemą, kuri atitinka Jungtinių Valstijų Maisto ir Vaistų administracijos **21 CFR 820 dalis**, **EN ISO 13485:2012**, Europos sąjungos tarybos direktyvą 93/42/EEB ir Japonijos PAL įstatymas.

Ballaigues,  
 2014 m. lapkričio 24 d.

Encl.



VERTIMAS TEISINGAS

UAB „Plandent“ direktorius Andrius Preišgalavičius



Maillefer Instruments Holding Sarl  
 Luc Etienne  
 Pardavimų direktorius

Produkto ir rūšies žymėjimas: <b>Priedai</b>		
Bylos numeris: <b>A35</b>		
Paruošė: Yassin Rouhila	Parašas:	Data: 2014-11-24
Peržiūrėjo: Luc ETIENNE	Parašas:	Data: 2014-11-24

**Dokumento istorija:**

Versija	Data	Pakeista santrauka
06	2014-11-24	Atnaujintas informacijos sąrašas.

Kodas	REF	Produkto pavadinimas	Paskirtis	GMDN	GMDN apibrėžimas	Tai-syklė	Klasė	S/M	Standartai
	A0197	Silikoninio stoperio dispenseris	Paprastas ir greitas būdas uždėti silikoninį stoperį ant šaknies kanalo instrument. Kai paspaudžiamas baltas mygtukas, stoperis pasirodo ir su instrument galiuku jį galima lengvai paimti.	38442	Neklasifikuojama	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041
	A0197A	Silikoninis stoperis	Priedas reguliuoti darbinį ilgį.	38442	Neklasifikuojama	5-1 hyphen	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1
R	A203M-1	Split-Kit M tuščias	Sterilizuojamas rinkinys kuris naudojamas laikyti instrumentams.	38442	Neklasifikuojama	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1
N	A203M-9	Split-Kit M tuščias	Sterilizuojamas rinkinys kuris naudojamas laikyti instrumentams.	38442	Neklasifikuojama	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1
	A203V	Split-Kit M tuščias	Sterilizuojamas rinkinys kuris naudojamas laikyti instrumentams.	38442	Neklasifikuojama	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1
D	A0207	Endo-kontaineris pilnas	Rinkinys naudojamas laikyti Gutaperčos, popierinius kaiščius ar kitus instrumentus.	38442	Neklasifikuojama	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1
D	A207T	Tūba Endo kontaineris	Tūba naudojamas laikyti Gutaperčos, popierinius kaiščius ar kitus instrumentus.	38442	Neklasifikuojama	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1
	A219P	Plastikinė mandrela	Rankiniam naudojimui šaknų kanalų instrumentų.	38442	Neklasifikuojama	5-1 hyphen	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1
R	A0227	Metalinė skėlimo dalis		38442	Neklasifikuojama	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1
R	A227T	Tūba	Gutaperčos, popierinių kaiščių ar kitų instrument laikymui.	38442	Neklasifikuojama	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041
	A259A	Cotton pellet	Prietaisas leidžiantis papildyti dezinfekcinio tirpalo.	38442	Neklasifikuojama	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041
	A265P	Endo stovelis	Sterilizuojamas rinkinys kuris naudojamas laikyti instrumentams.	38442	Neklasifikuojama	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1
	A0266	Endo stovelis (organizier) tuščias	Sterilizuojamas rinkinys ProFile sistemai.	38442	Neklasifikuojama	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1
	A268P	Clean (kempinėlių) stovelis	Naudojamas laikyti instrumentus švarioje vietoje endodontinių procedure metu.	38442	Neklasifikuojama	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1
	A268P-2	Didelis Clean (kempinėlių) stovelis	Naudojamas laikyti instrumentus švarioje vietoje endodontinių procedure metu.	38442	Neklasifikuojama	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1
D	A268S	Clean (kempinėlių) stovelio kempinėlis	Clean (kempinėlių) stovelio putos blokeliai Ref. A268P	38442	Neklasifikuojama	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1
D	A268S-2	Didelio Clean (kempinėlių) stovelio kempinėlis	Clean (kempinėlių) stovelio putos blokeliai Ref. A268P-2	38442	Neklasifikuojama	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1
	A272V	Sosio dėžutė tuščia	Sterilizuojamas rinkinys kuris naudojamas laikyti instrumentams.	38442	Neklasifikuojama	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1
	A274V	Flexokit tuščias	Sterilizuojamas rinkinys kuris naudojamas laikyti instrumentams.	38442	Neklasifikuojama	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1
	A0290	Įdėklas endo Prof.Bogopolsky	Plastikinis įdėklas su skylutėmis instrumentams ir putos bloku galima naudoti Split rinkinyje.	38442	Neklasifikuojama	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1
R	A0349	Multi stovelis	Sterilizuojamas rinkinys ProFile sistemai.	38442	Neklasifikuojama	1	I	-	ISO 13485, ISO 14971, ISO 15223-1, EN 1041, ISO 17664, ISO 17665-1
	A0362-9	ProFile stovelis	Sterilizuojamas rinkinys ProFile sistemai.	38442	Neklasifikuojama	1	I	-	ISO 13485, ISO 14971, ISO



*Document Detail*

**Type:** 1450-RDT\_Dec of Conf  
**Document No.:** 1450-RDT\_Dec of Conf\_B01[2]  
**Title:** B01-Declaration of conformity  
**Comment** B01-Declaration of conformity 02  
**Status:** CURRENT  
**Effective Date:** 10-Jan-2009

## CE declaration of conformity – Directive 93/42/EEC

**Class I products**

We

**MAILLEFER INSTRUMENTS HOLDING Sàrl and its subsidiaries**

**Chemin du Verger 3, CH-1338 Ballaigues / Switzerland**

declare on our own responsibility that the following products listed in enclosure F 711.208.E:  
**Reference List B01** (1 page), version n°02.

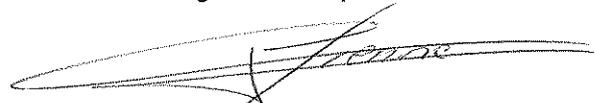
are conform to the essential requirements and provisions of Council Directive 93/42/EEC.

This declaration is based on the following data:

- Technical documentation demonstrating the product conformity to the requirements of the directive and the edict.
- Applied norms: Listed in enclosure F 711.208.E: **Reference List B01**
- Conformity assessment procedure: Annex VII of the directive 93/42/EEC

Ballaigues, 01.10.2009

Management Representative



Luc Etienne

MAILLEFER INSTRUMENTS  
HOLDING Sàrl

Encl.

B01: Hand instruments

Code: D : Displacement from other list / N : New reference / M : Modification / R : Removed  
 GMDN : Global Medical Device Nomenclature  
 REF : Maillefer reference number  
 S : sterile / M : measure

Edition : 02

Date: 01.10.2009

Signature :



Code	REF	Proprietary Name	GMDN #	GMDN definition	Rule	Class	S/M	Standards
	B0093	Probe simple or double ended	31848	Probe, dental	6-2nd hyphen	I	-	EN ISO 13485, ISO 14971, EN 1639, ISO 7492 (part.), ISO 17664, ISO 17665-1, EN 980, ISO 15223-1, EN 1041
	B093S	Probe simple or double ended	31848	Probe, dental	6-2nd hyphen	I	-	EN ISO 13485, ISO 14971, EN 1639, ISO 7492 (part.), ISO 17664, ISO 17665-1, EN 980, ISO 15223-1, EN 1041
	B0095	Excavators double ended	35811	Excavator, dental, hand held	5-1st hyphen	I	-	EN ISO 13485, ISO 14971, EN 1639, ISO 7492 (part.), ISO 17664, ISO 17665-1, EN 980, ISO 15223-1, EN 1041
	B095S	Excavator double ended	35811	Excavator, dental, hand held	5-1st hyphen	I	-	EN ISO 13485, ISO 14971, EN 1639, ISO 7492 (part.), ISO 17664, ISO 17665-1, EN 980, ISO 15223-1, EN 1041
	B0188	Mouth mirror handle	31776	Mirror, dental, hand held	5-1st hyphen	I	-	EN ISO 13485, ISO 14971, EN 1639, ISO 7492 (part.), ISO 17664, ISO 17665-1, EN 980, ISO 15223-1, EN 1041



**ATTESTATION / CERTIFICATE / BESCHEINIGUNG n° 10468 rev. 4**

**Délivrée à Paris le 20 août 2015**

*Issued in Paris on August 20th, 2015*

*Ausgestellt in Paris, den 20 August 2015*

## **ATTESTATION CE / EC CERTIFICATE / EG-BESCHEINIGUNG**

**Approbation du Système Complet d'assurance Qualité / Approval of full Quality Assurance System**

**Genehmigung von dem vollständiges Qualitätssicherungssystem**

**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*

*Anhang II, ohne den punkt 4, Richtlinie 93/42/EWG über Medizinprodukte*

**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 / Hersteller**

**MAILLEFER INSTRUMENTS HOLDING SARL**

**Chemin du Verger 3,**

**1338 BALLAIGUES SWITZERLAND**

**Catégorie du(des) dispositif(s) / Device(s) category / Produkt-Kategorie**

**Instruments dentaires stériles et non stériles, moteurs et contre-angles, système de reconstruction dentaire, appareils destinés au diagnostic dans l'art dentaire**

*Sterile and non sterile dental instruments, motors and contra angles, dental reconstruction systems, dental devices for diagnoses*

*Sterile und nicht sterile Dentalinstrumente, Motoren and Winkelstücke, Dentalwiederaufbausysteme, Geräte für Diagnose im Dentalbereich*

*Voir détails sur addendum / See attachment for additional information*

**Le LNE/G-MED atteste qu'à l'examen des résultats figurant dans le rapport référencé P129078, 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.**

*LNE/G-MED certifies that, on the basis of the results contained in the file referenced P129078, 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.*

*LNE/G-MED bescheinigt, dass nach Überprüfung der im Prüfbericht n° P129078, enthaltenen Ergebnisse, das Qualitätssicherungssystem – für Designlenkung, Herstellung und Endkontrolle – der o.g. aufgeführten Medizinprodukte, den Anforderungen der Anhang II ohne den punkt 4 der Richtlinie 93/42/EWG entspricht..*

**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 / Gültig ab : August 20th, 2015 (included)**

**Valable jusqu'au / Expiry date / Gültig bis : July 8th, 2018 (included)**



**On behalf of the Certification Director**

**Cécile VAUGÉADE**

**G-MED Certification Division Manager**

**der stellvertretende Direktor i.V.**

### Identification des dispositifs / Identification of devices

Les produits couverts par ce certificat sont référencés sur la liste des produits (55 pages) authentifiée par le LNE/G-MED en date du 20 août 2015 / *The products covered by this certificate are listed on the manufacturer's list of products (55 pages) authenticated by LNE/G-MED on August 20, 2015.*

### Identification des sites et activités / Identification of locations and activities

- **Chemin du Verger, 3, 1338 BALLAIGUES, SUISSE :**  
Conception, fabrication, contrôle final / *Design, manufacturing, final control*
- **Chemin de Beau-Site, 5, 1338 BALLAIGUES, SUISSE :**  
Fabrication / *Manufacturing*
- **Le Creux, 1338 BALLAIGUES, SUISSE :**  
Fabrication / *Manufacturing*

LNE/G-MED

0459



On behalf of the Certification Director  
**Cécile VAUGELADE**  
G-MED Certification Division Manager

**Liste des produits couverts par l'attestation CE n°10468 rev.4**  
**List of products covered by the EC Certificate n°10468 rev.4**  
**Liste der produkte, welch emit dem EG Zertifikat n°10468 rev.4 abgedeckt werden**

Item number	Product designation	Product class
60667332	X-Post	Class IIa
60667333	X-Post	Class IIa
60667334	X-Post	Class IIa
60667335	X-Post	Class IIa
A000724000100	Root canal drill	Class IIa
A000724000200	Root canal drill	Class IIa
A000724000300	Root canal drill	Class IIa
A000724000400	Root canal drill	Class IIa
A000724090000	Root canal drill	Class IIa
A000823000100	Gates glidden drill	Class IIa
A000823000103	Gates sterile glidden drill	Class IIa
A000823000200	Gates glidden drill	Class IIa
A000823000203	Gates sterile glidden drill	Class IIa
A000823000300	Gates glidden drill	Class IIa
A000823000303	Gates sterile glidden drill	Class IIa
A000823000400	Gates glidden drill	Class IIa
A000823000403	Gates sterile glidden drill	Class IIa
A000823000500	Gates glidden drill	Class IIa
A000823000503	Gates sterile glidden drill	Class IIa
A000823000600	Gates glidden drill	Class IIa
A000823000603	Gates sterile glidden drill	Class IIa
A000823090100	Gates glidden drill	Class IIa
A000823090103	Gates sterile glidden drill	Class IIa
A000824000100	Gates glidden drill	Class IIa
A000824000103	Gates sterile glidden drill	Class IIa
A000824000200	Gates glidden drill	Class IIa
A000824000203	Gates sterile glidden drill	Class IIa
A000824000300	Gates glidden drill	Class IIa
A000824000303	Gates sterile glidden drill	Class IIa
A000824000400	Gates glidden drill	Class IIa
A000824000403	Gates sterile glidden drill	Class IIa
A000824000500	Gates glidden drill	Class IIa
A000824000503	Gates sterile glidden drill	Class IIa
A000824000600	Gates glidden drill	Class IIa
A000824000603	Gates sterile glidden drill	Class IIa
A000824000700	Gates glidden drill	Class IIa
A000824000703	Gates sterile glidden drill	Class IIa
A000824090100	Gates glidden drill	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits

LNE/G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed

20-AUG-2015



Item number	Product designation	Product class
A000824090103	Gates sterile glidden drill	Class IIa
A000825000100	Gates glidden drill	Class IIa
A000825000103	Gates sterile glidden drill	Class IIa
A000825000200	Gates glidden drill	Class IIa
A000825000203	Gates sterile glidden drill	Class IIa
A000825000300	Gates glidden drill	Class IIa
A000825000303	Gates sterile glidden drill	Class IIa
A000825000400	Gates glidden drill	Class IIa
A000825000403	Gates sterile glidden drill	Class IIa
A000825000500	Gates glidden drill	Class IIa
A000825000503	Gates sterile glidden drill	Class IIa
A000825000600	Gates glidden drill	Class IIa
A000825090100	Gates glidden drill	Class IIa
A000825090103	Gates glidden drill	Class IIa
A000923000100	Largo Peeso Reamer	Class IIa
A000923000103	Largo Sterile	Class IIa
A000923000200	Largo Peeso Reamer	Class IIa
A000923000203	Largo Sterile	Class IIa
A000923000300	Largo Peeso Reamer	Class IIa
A000923000303	Largo Sterile	Class IIa
A000923000400	Largo Peeso Reamer	Class IIa
A000923000403	Largo Sterile	Class IIa
A000923000500	Largo Peeso Reamer	Class IIa
A000923000503	Largo Sterile	Class IIa
A000923000600	Largo Peeso Reamer	Class IIa
A000923000603	Largo Sterile	Class IIa
A000923090100	Largo Peeso Reamer	Class IIa
A000923090103	Largo Sterile	Class IIa
A000924000100	Largo Peeso Reamer	Class IIa
A000924000103	Largo Sterile	Class IIa
A000924000200	Largo Peeso Reamer	Class IIa
A000924000203	Largo Sterile	Class IIa
A000924000300	Largo Peeso Reamer	Class IIa
A000924000303	Largo Sterile	Class IIa
A000924000400	Largo Peeso Reamer	Class IIa
A000924000403	Largo Sterile	Class IIa
A000924000500	Largo Peeso Reamer	Class IIa
A000924000503	Largo Sterile	Class IIa
A000924000600	Largo Peeso Reamer	Class IIa
A000924000603	Largo Sterile	Class IIa
A000924090100	Largo Peeso Reamer	Class IIa
A000924090103	Largo Sterile	Class IIa
A001012501000	K-Reamer Torpan	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits  
 LNE/G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed

Item number	Product designation	Product class
A001012501500	K-Reamer Torpan	Class IIa
A001012502000	K-Reamer Torpan	Class IIa
A001012502500	K-Reamer Torpan	Class IIa
A001012503000	K-Reamer Torpan	Class IIa
A001012503500	K-Reamer Torpan	Class IIa
A001012504000	K-Reamer Torpan	Class IIa
A001012504500	K-Reamer Torpan	Class IIa
A001012505000	K-Reamer Torpan	Class IIa
A001012505500	K-Reamer Torpan	Class IIa
A001012506000	K-Reamer Torpan	Class IIa
A001012507000	K-Reamer Torpan	Class IIa
A001012508000	K-Reamer Torpan	Class IIa
A001012509000	K-Reamer Torpan	Class IIa
A001012510000	K-Reamer Torpan	Class IIa
A001012512000	K-Reamer Torpan	Class IIa
A001012590000	K-Reamer Torpan	Class IIa
A001012590100	K-Reamer Torpan	Class IIa
A001021801000	K-Reamer Torpan	Class IIa
A001021801500	K-Reamer Torpan	Class IIa
A001021802000	K-Reamer Torpan	Class IIa
A001021802500	K-Reamer Torpan	Class IIa
A001021803000	K-Reamer Torpan	Class IIa
A001021803500	K-Reamer Torpan	Class IIa
A001021804000	K-Reamer Torpan	Class IIa
A001021804500	K-Reamer Torpan	Class IIa
A001021805000	K-Reamer Torpan	Class IIa
A001021805500	K-Reamer Torpan	Class IIa
A001021806000	K-Reamer Torpan	Class IIa
A001021807000	K-Reamer Torpan	Class IIa
A001021808000	K-Reamer Torpan	Class IIa
A001021809000	K-Reamer Torpan	Class IIa
A001021810000	K-Reamer Torpan	Class IIa
A001021811000	K-Reamer Torpan	Class IIa
A001021812000	K-Reamer Torpan	Class IIa
A001021890000	K-Reamer Torpan	Class IIa
A001021890100	K-Reamer Torpan	Class IIa
A001022101000	K-Reamer Torpan	Class IIa
A001022101500	K-Reamer Torpan	Class IIa
A001022102000	K-Reamer Torpan	Class IIa
A001022102500	K-Reamer Torpan	Class IIa
A001022103000	K-Reamer Torpan	Class IIa
A001022103500	K-Reamer Torpan	Class IIa
A001022104000	K-Reamer Torpan	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est  
 valide pour les dispositifs médicaux décrits  
 LNE/G-MED (0459) recognizes that its EC certificate  
 is valid for the medical devices listed

Item number	Product designation	Product class
A001022104500	K-Reamer Torpan	Class IIa
A001022105000	K-Reamer Torpan	Class IIa
A001022105500	K-Reamer Torpan	Class IIa
A001022106000	K-Reamer Torpan	Class IIa
A001022107000	K-Reamer Torpan	Class IIa
A001022108000	K-Reamer Torpan	Class IIa
A001022109000	K-Reamer Torpan	Class IIa
A001022110000	K-Reamer Torpan	Class IIa
A001022111000	K-Reamer Torpan	Class IIa
A001022112000	K-Reamer Torpan	Class IIa
A001022190000	K-Reamer Torpan	Class IIa
A001022190100	K-Reamer Torpan	Class IIa
A001022501000	K-Reamer Torpan	Class IIa
A001022501500	K-Reamer Torpan	Class IIa
A001022502000	K-Reamer Torpan	Class IIa
A001022502500	K-Reamer Torpan	Class IIa
A001022503000	K-Reamer Torpan	Class IIa
A001022503500	K-Reamer Torpan	Class IIa
A001022504000	K-Reamer Torpan	Class IIa
A001022504500	K-Reamer Torpan	Class IIa
A001022505000	K-Reamer Torpan	Class IIa
A001022505500	K-Reamer Torpan	Class IIa
A001022506000	K-Reamer Torpan	Class IIa
A001022506003	K-Reamer Torpan	Class IIa
A001022507000	K-Reamer Torpan	Class IIa
A001022508000	K-Reamer Torpan	Class IIa
A001022509000	K-Reamer Torpan	Class IIa
A001022510000	K-Reamer Torpan	Class IIa
A001022511000	K-Reamer Torpan	Class IIa
A001022512000	K-Reamer Torpan	Class IIa
A001022590000	K-Reamer Torpan	Class IIa
A001022590100	K-Reamer Torpan	Class IIa
A001522101300	PATHFILE	Class IIa
A001522101303	PATHFILE STERILE	Class IIa
A001522101600	PATHFILE	Class IIa
A001522101603	PATHFILE STERILE	Class IIa
A001522101900	PATHFILE	Class IIa
A001522101903	PATHFILE STERILE	Class IIa
A001522190000	PATHFILE	Class IIa
A001522190003	PATHFILE STERILE	Class IIa
A001522501300	PATHFILE	Class IIa
A001522501303	PATHFILE STERILE	Class IIa
A001522501600	PATHFILE	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits

LNE/G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed

Item number	Product designation	Product class
A001522501603	PATHFILE STERILE	Class IIa
A001522501900	PATHFILE	Class IIa
A001522501903	PATHFILE STERILE	Class IIa
A001522590000	PATHFILE	Class IIa
A001522590003	PATHFILE STERILE	Class IIa
A001522590200	PATHFILE	Class IIa
A001523101300	PATHFILE	Class IIa
A001523101303	PATHFILE STERILE	Class IIa
A001523101600	PATHFILE	Class IIa
A001523101603	PATHFILE STERILE	Class IIa
A001523101900	PATHFILE	Class IIa
A001523101903	PATHFILE STERILE	Class IIa
A001523190000	PATHFILE	Class IIa
A001523190003	PATHFILE STERILE	Class IIa
A002220090000	Lentulo Past Carrier	Class IIa
A002221700100	Lentulo Past Carrier	Class IIa
A002221700103	Lentulo sterile Paste Carrier	Class IIa
A002221700200	Lentulo Past Carrier	Class IIa
A002221700203	Lentulo sterile Paste Carrier	Class IIa
A002221700300	Lentulo Past Carrier	Class IIa
A002221700303	Lentulo sterile Paste Carrier	Class IIa
A002221700400	Lentulo Past Carrier	Class IIa
A002221700403	Lentulo sterile Paste Carrier	Class IIa
A002221790000	Lentulo Past Carrier	Class IIa
A002221790100	Lentulo Past Carrier	Class IIa
A002222100100	Lentulo Past Carrier	Class IIa
A002222100103	Lentulo sterile Paste Carrier	Class IIa
A002222100200	Lentulo Past Carrier	Class IIa
A002222100203	Lentulo sterile Paste Carrier	Class IIa
A002222100300	Lentulo Past Carrier	Class IIa
A002222100303	Lentulo sterile Paste Carrier	Class IIa
A002222100400	Lentulo Past Carrier	Class IIa
A002222100403	Lentulo sterile Paste Carrier	Class IIa
A002222190000	Lentulo Past Carrier	Class IIa
A002222190003	Lentulo sterile Paste Carrier	Class IIa
A002222190100	Lentulo Past Carrier	Class IIa
A002222500100	Lentulo Past Carrier	Class IIa
A002222500103	Lentulo sterile Paste Carrier	Class IIa
A002222500200	Lentulo Past Carrier	Class IIa
A002222500203	Lentulo sterile Paste Carrier	Class IIa
A002222500300	Lentulo Past Carrier	Class IIa
A002222500303	Lentulo sterile Paste Carrier	Class IIa
A002222500400	Lentulo Past Carrier	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est  
 valide pour les dispositifs médicaux décrits  
 ..... 2015  
 LNE/G-MED (0459) recognizes that its EC certificate  
 is valid for the medical devices listed

Item number	Product designation	Product class
A002222500403	Lentulo sterile Paste Carrier	Class IIa
A002222590000	Lentulo Past Carrier	Class IIa
A002222590003	Lentulo sterile Paste Carrier	Class IIa
A002222590100	Lentulo Past Carrier	Class IIa
A004B02102001	Spiro COLORINOX	Class IIa
A004B02102501	Spiro COLORINOX	Class IIa
A004B02103001	Spiro COLORINOX	Class IIa
A004B02103501	Spiro COLORINOX	Class IIa
A004B02104001	Spiro COLORINOX	Class IIa
A004B02105001	Spiro COLORINOX	Class IIa
A004B02106001	Spiro COLORINOX	Class IIa
A004B02190001	Spiro COLORINOX	Class IIa
A004C02102001	Spiro Short Barbed Nervbroach	Class IIa
A004C02102501	Spiro Short Barbed Nervbroach	Class IIa
A004C02103001	Spiro Short Barbed Nervbroach	Class IIa
A004C02103501	Spiro Short Barbed Nervbroach	Class IIa
A004C02104001	Spiro Short Barbed Nervbroach	Class IIa
A004C02105001	Spiro Short Barbed Nervbroach	Class IIa
A004C02106001	Spiro Short Barbed Nervbroach	Class IIa
A005032501000	THERMA-CUT	Class IIa
A005032501200	THERMA-CUT	Class IIa
A005032501400	THERMA-CUT	Class IIa
A005032501600	THERMA-CUT	Class IIa
A005032590000	THERMA-CUT	Class IIa
A005131300500	POST SPACE BUR	Class IIa
A005131300700	POST SPACE BUR	Class IIa
A005131900500	POST SPACE BUR	Class IIa
A005131900700	POST SPACE BUR	Class IIa
A011C02101003	K-Flexoreamer sterile	Class IIa
A011C02101503	K-Flexoreamer sterile	Class IIa
A011C02102003	K-Flexoreamer sterile	Class IIa
A011C02102503	K-Flexoreamer sterile	Class IIa
A011C02103003	K-Flexoreamer sterile	Class IIa
A011C02103503	K-Flexoreamer sterile	Class IIa
A011C02104003	K-Flexoreamer sterile	Class IIa
A011C02104503	K-Flexoreamer sterile	Class IIa
A011C02105003	K-Flexoreamer sterile	Class IIa
A011C02105503	K-Flexoreamer sterile	Class IIa
A011C02106003	K-Flexoreamer sterile	Class IIa
A011C02107003	K-Flexoreamer sterile	Class IIa
A011C02108003	K-Flexoreamer sterile	Class IIa
A011C02190003	K-Flexoreamer sterile	Class IIa
A011C02190103	K-Flexoreamer sterile	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits  
 LNE/G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed

Item number	Product designation	Product class
A011C02190203	K-Flexoreamer sterile	Class IIa
A011C02500803	K-Flexoreamer sterile	Class IIa
A011C02501003	K-Flexoreamer sterile	Class IIa
A011C02501503	K-Flexoreamer sterile	Class IIa
A011C02502003	K-Flexoreamer sterile	Class IIa
A011C02502503	K-Flexoreamer sterile	Class IIa
A011C02503003	K-Flexoreamer sterile	Class IIa
A011C02503503	K-Flexoreamer sterile	Class IIa
A011C02504003	K-Flexoreamer sterile	Class IIa
A011C02504503	K-Flexoreamer sterile	Class IIa
A011C02505003	K-Flexoreamer sterile	Class IIa
A011C02505503	K-Flexoreamer sterile	Class IIa
A011C02506003	K-Flexoreamer sterile	Class IIa
A011C02507003	K-Flexoreamer sterile	Class IIa
A011C02508003	K-Flexoreamer sterile	Class IIa
A011C02590003	K-Flexoreamer sterile	Class IIa
A011C02590103	K-Flexoreamer sterile	Class IIa
A011C02590203	K-Flexoreamer sterile	Class IIa
A011C03101503	K-Flexoreamer sterile	Class IIa
A011C03102003	K-Flexoreamer sterile	Class IIa
A011C03102503	K-Flexoreamer sterile	Class IIa
A011C03103003	K-Flexoreamer sterile	Class IIa
A011C03103503	K-Flexoreamer sterile	Class IIa
A011C03104003	K-Flexoreamer sterile	Class IIa
A011C03190003	K-Flexoreamer sterile	Class IIa
A011N21801500	PROFILE.04	Class IIa
A011N21801503	PROFILE STERILE .04	Class IIa
A011N21802000	PROFILE.04	Class IIa
A011N21802003	PROFILE STERILE .04	Class IIa
A011N21802500	PROFILE.04	Class IIa
A011N21802503	PROFILE STERILE .04	Class IIa
A011N21803000	PROFILE.04	Class IIa
A011N21803003	PROFILE STERILE .04	Class IIa
A011N21803500	PROFILE.04	Class IIa
A011N21803503	PROFILE STERILE .04	Class IIa
A011N21804000	PROFILE.04	Class IIa
A011N21804003	PROFILE STERILE .04	Class IIa
A011N21890000	PROFILE.04	Class IIa
A011N21890003	PROFILE STERILE .04	Class IIa
A011N22101500	PROFILE.04	Class IIa
A011N22101503	PROFILE STERILE .04	Class IIa
A011N22102000	PROFILE.04	Class IIa
A011N22102003	PROFILE STERILE .04	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits

LNE/G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed

28 AOÛT 2015

Item number	Product designation	Product class
A011N22102500	PROFILE.04	Class IIa
A011N22102503	PROFILE STERILE .04	Class IIa
A011N22103000	PROFILE.04	Class IIa
A011N22103003	PROFILE STERILE .04	Class IIa
A011N22103500	PROFILE.04	Class IIa
A011N22103503	PROFILE STERILE .04	Class IIa
A011N22104000	PROFILE.04	Class IIa
A011N22104003	PROFILE STERILE .04	Class IIa
A011N22104500	PROFILE.04	Class IIa
A011N22104503	PROFILE STERILE .04	Class IIa
A011N22106000	PROFILE.04	Class IIa
A011N22106003	PROFILE STERILE .04	Class IIa
A011N22109000	PROFILE.04	Class IIa
A011N22109003	PROFILE STERILE .04	Class IIa
A011N22190000	PROFILE.04	Class IIa
A011N22190003	PROFILE STERILE .04	Class IIa
A011N22190700	PROFILE.04	Class IIa
A011N22190703	PROFILE STERILE .04	Class IIa
A011N22501500	PROFILE.04	Class IIa
A011N22501503	PROFILE STERILE .04	Class IIa
A011N22502000	PROFILE.04	Class IIa
A011N22502003	PROFILE STERILE .04	Class IIa
A011N22502500	PROFILE.04	Class IIa
A011N22502503	PROFILE STERILE .04	Class IIa
A011N22503000	PROFILE.04	Class IIa
A011N22503003	PROFILE STERILE .04	Class IIa
A011N22503500	PROFILE.04	Class IIa
A011N22503503	PROFILE STERILE .04	Class IIa
A011N22504000	PROFILE.04	Class IIa
A011N22504003	PROFILE STERILE .04	Class IIa
A011N22504500	PROFILE.04	Class IIa
A011N22504503	PROFILE STERILE .04	Class IIa
A011N22506000	PROFILE.04	Class IIa
A011N22506003	PROFILE STERILE .04	Class IIa
A011N22509000	PROFILE.04	Class IIa
A011N22509003	PROFILE STERILE .04	Class IIa
A011N22590000	PROFILE.04	Class IIa
A011N22590003	PROFILE STERILE .04	Class IIa
A011N22590700	PROFILE.04	Class IIa
A011N22590703	PROFILE STERILE .04	Class IIa
A011N23101500	PROFILE.04	Class IIa
A011N23101503	PROFILE STERILE .04	Class IIa
A011N23102000	PROFILE.04	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits ..... 2015  
 LNE/G-MED (0458) recognizes that its EC certificate is valid for the medical devices listed

Item number	Product designation	Product class
A011N23102003	PROFILE STERILE .04	Class IIa
A011N23102500	PROFILE.04	Class IIa
A011N23102503	PROFILE STERILE .04	Class IIa
A011N23103000	PROFILE.04	Class IIa
A011N23103003	PROFILE STERILE .04	Class IIa
A011N23103500	PROFILE.04	Class IIa
A011N23103503	PROFILE STERILE .04	Class IIa
A011N23104000	PROFILE.04	Class IIa
A011N23104003	PROFILE STERILE .04	Class IIa
A011N23104500	PROFILE.04	Class IIa
A011N23104503	PROFILE STERILE .04	Class IIa
A011N23106000	PROFILE.04	Class IIa
A011N23106003	PROFILE STERILE .04	Class IIa
A011N23109000	PROFILE.04	Class IIa
A011N23109003	PROFILE STERILE .04	Class IIa
A011N23190000	PROFILE.04	Class IIa
A011N23190003	PROFILE STERILE .04	Class IIa
A011N23190700	PROFILE.04	Class IIa
A011N23190703	PROFILE STERILE .04	Class IIa
A012C01801503	K-Flexofile Sterile	Class IIa
A012C01802003	K-Flexofile Sterile	Class IIa
A012C01802503	K-Flexofile Sterile	Class IIa
A012C01803003	K-Flexofile Sterile	Class IIa
A012C01803503	K-Flexofile Sterile	Class IIa
A012C01804003	K-Flexofile Sterile	Class IIa
A012C01890003	K-Flexofile Sterile	Class IIa
A012C02100603	K-Flexofile Sterile	Class IIa
A012C02100803	K-Flexofile Sterile	Class IIa
A012C02101003	K-Flexofile Sterile	Class IIa
A012C02101503	K-Flexofile Sterile	Class IIa
A012C02102003	K-Flexofile Sterile	Class IIa
A012C02102503	K-Flexofile Sterile	Class IIa
A012C02103003	K-Flexofile Sterile	Class IIa
A012C02103503	K-Flexofile Sterile	Class IIa
A012C02104003	K-Flexofile Sterile	Class IIa
A012C02104503	K-Flexofile Sterile	Class IIa
A012C02105003	K-Flexofile Sterile	Class IIa
A012C02105503	K-Flexofile Sterile	Class IIa
A012C02106003	K-Flexofile Sterile	Class IIa
A012C02107003	K-Flexofile Sterile	Class IIa
A012C02108003	K-Flexofile Sterile	Class IIa
A012C02109003	K-Flexofile Sterile	Class IIa
A012C02110003	K-Flexofile Sterile	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits  
 LNE/G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed  
 21.08.2015

Item number	Product designation	Product class
A012C02111003	K-Flexofile Sterile	Class IIa
A012C02112003	K-Flexofile Sterile	Class IIa
A012C02113003	K-Flexofile Sterile	Class IIa
A012C02114003	K-Flexofile Sterile	Class IIa
A012C02190003	K-Flexofile Sterile	Class IIa
A012C02190103	K-Flexofile Sterile	Class IIa
A012C02190203	K-Flexofile Sterile	Class IIa
A012C02500603	K-Flexofile Sterile	Class IIa
A012C02500803	K-Flexofile Sterile	Class IIa
A012C02501003	K-Flexofile Sterile	Class IIa
A012C02501503	K-Flexofile Sterile	Class IIa
A012C02502003	K-Flexofile Sterile	Class IIa
A012C02502503	K-Flexofile Sterile	Class IIa
A012C02503003	K-Flexofile Sterile	Class IIa
A012C02503503	K-Flexofile Sterile	Class IIa
A012C02504003	K-Flexofile Sterile	Class IIa
A012C02504503	K-Flexofile Sterile	Class IIa
A012C02505003	K-Flexofile Sterile	Class IIa
A012C02505503	K-Flexofile Sterile	Class IIa
A012C02506003	K-Flexofile Sterile	Class IIa
A012C02507003	K-Flexofile Sterile	Class IIa
A012C02508003	K-Flexofile Sterile	Class IIa
A012C02509003	K-Flexofile Sterile	Class IIa
A012C02510003	K-Flexofile Sterile	Class IIa
A012C02511003	K-Flexofile Sterile	Class IIa
A012C02512003	K-Flexofile Sterile	Class IIa
A012C02513003	K-Flexofile Sterile	Class IIa
A012C02514003	K-Flexofile Sterile	Class IIa
A012C02590003	K-Flexofile Sterile	Class IIa
A012C02590103	K-Flexofile Sterile	Class IIa
A012C02590203	K-Flexofile Sterile	Class IIa
A012C03100603	K-Flexofile Sterile	Class IIa
A012C03100803	K-Flexofile Sterile	Class IIa
A012C03101003	K-Flexofile Sterile	Class IIa
A012C03101503	K-Flexofile Sterile	Class IIa
A012C03102003	K-Flexofile Sterile	Class IIa
A012C03102503	K-Flexofile Sterile	Class IIa
A012C03103003	K-Flexofile Sterile	Class IIa
A012C03103503	K-Flexofile Sterile	Class IIa
A012C03104003	K-Flexofile Sterile	Class IIa
A012C03104503	K-Flexofile Sterile	Class IIa
A012C03105003	K-Flexofile Sterile	Class IIa
A012C03105503	K-Flexofile Sterile	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits  
 LNE/G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed

Item number	Product designation	Product class
A012C03106003	K-Flexofile Sterile	Class IIa
A012C03107003	K-Flexofile Sterile	Class IIa
A012C03108003	K-Flexofile Sterile	Class IIa
A012C03109003	K-Flexofile Sterile	Class IIa
A012C03110003	K-Flexofile Sterile	Class IIa
A012C03111003	K-Flexofile Sterile	Class IIa
A012C03112003	K-Flexofile Sterile	Class IIa
A012C03113003	K-Flexofile Sterile	Class IIa
A012C03114003	K-Flexofile Sterile	Class IIa
A012C03190003	K-Flexofile Sterile	Class IIa
A012C03190103	K-Flexofile Sterile	Class IIa
A012C03190203	K-Flexofile Sterile	Class IIa
A012D01800603	K-File COLORINOX sterile	Class IIa
A012D01800803	K-File COLORINOX sterile	Class IIa
A012D01801003	K-File COLORINOX sterile	Class IIa
A012D02100603	K-File COLORINOX sterile	Class IIa
A012D02100803	K-File COLORINOX sterile	Class IIa
A012D02101003	K-File COLORINOX sterile	Class IIa
A012D02101503	K-File COLORINOX sterile	Class IIa
A012D02102003	K-File COLORINOX sterile	Class IIa
A012D02102503	K-File COLORINOX sterile	Class IIa
A012D02103003	K-File COLORINOX sterile	Class IIa
A012D02103503	K-File COLORINOX sterile	Class IIa
A012D02104003	K-File COLORINOX sterile	Class IIa
A012D02104503	K-File COLORINOX sterile	Class IIa
A012D02105003	K-File COLORINOX sterile	Class IIa
A012D02105503	K-File COLORINOX sterile	Class IIa
A012D02106003	K-File COLORINOX sterile	Class IIa
A012D02107003	K-File COLORINOX sterile	Class IIa
A012D02108003	K-File COLORINOX sterile	Class IIa
A012D02109003	K-File COLORINOX sterile	Class IIa
A012D02110003	K-File COLORINOX sterile	Class IIa
A012D02111003	K-File COLORINOX sterile	Class IIa
A012D02112003	K-File COLORINOX sterile	Class IIa
A012D02113003	K-File COLORINOX sterile	Class IIa
A012D02114003	K-File COLORINOX sterile	Class IIa
A012D02190003	K-File COLORINOX sterile	Class IIa
A012D02190103	K-File COLORINOX sterile	Class IIa
A012D02190203	K-File COLORINOX sterile	Class IIa
A012D02500603	K-File COLORINOX sterile	Class IIa
A012D02500803	K-File COLORINOX sterile	Class IIa
A012D02501003	K-File COLORINOX sterile	Class IIa
A012D02501503	K-File COLORINOX sterile	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits ..... 20 AOUT 2015  
 LNE/G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed

Item number	Product designation	Product class
A012D02502003	K-File COLORINOX sterile	Class IIa
A012D02502503	K-File COLORINOX sterile	Class IIa
A012D02503003	K-File COLORINOX sterile	Class IIa
A012D02503503	K-File COLORINOX sterile	Class IIa
A012D02504003	K-File COLORINOX sterile	Class IIa
A012D02504503	K-File COLORINOX sterile	Class IIa
A012D02505003	K-File COLORINOX sterile	Class IIa
A012D02505503	K-File COLORINOX sterile	Class IIa
A012D02506003	K-File COLORINOX sterile	Class IIa
A012D02507003	K-File COLORINOX sterile	Class IIa
A012D02508003	K-File COLORINOX sterile	Class IIa
A012D02509003	K-File COLORINOX sterile	Class IIa
A012D02510003	K-File COLORINOX sterile	Class IIa
A012D02511003	K-File COLORINOX sterile	Class IIa
A012D02512003	K-File COLORINOX sterile	Class IIa
A012D02513003	K-File COLORINOX sterile	Class IIa
A012D02514003	K-File COLORINOX sterile	Class IIa
A012D02590003	K-File COLORINOX sterile	Class IIa
A012D02590103	K-File COLORINOX sterile	Class IIa
A012D02590203	K-File COLORINOX sterile	Class IIa
A012D02800603	K-File COLORINOX sterile	Class IIa
A012D02800803	K-File COLORINOX sterile	Class IIa
A012D02801003	K-File COLORINOX sterile	Class IIa
A012D02801503	K-File COLORINOX sterile	Class IIa
A012D02802003	K-File COLORINOX sterile	Class IIa
A012D02802503	K-File COLORINOX sterile	Class IIa
A012D02803003	K-File COLORINOX sterile	Class IIa
A012D02803503	K-File COLORINOX sterile	Class IIa
A012D02804003	K-File COLORINOX sterile	Class IIa
A012D02804503	K-File COLORINOX sterile	Class IIa
A012D02805003	K-File COLORINOX sterile	Class IIa
A012D02805503	K-File COLORINOX sterile	Class IIa
A012D02806003	K-File COLORINOX sterile	Class IIa
A012D02807003	K-File COLORINOX sterile	Class IIa
A012D02808003	K-File COLORINOX sterile	Class IIa
A012D02809003	K-File COLORINOX sterile	Class IIa
A012D02810003	K-File COLORINOX sterile	Class IIa
A012D02811003	K-File COLORINOX sterile	Class IIa
A012D02812003	K-File COLORINOX sterile	Class IIa
A012D02813003	K-File COLORINOX sterile	Class IIa
A012D02814003	K-File COLORINOX sterile	Class IIa
A012D02890003	K-File COLORINOX sterile	Class IIa
A012D02890103	K-File COLORINOX sterile	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits ..... 20.01.2016  
 LNE/G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed

Item number	Product designation	Product class
A012D02890203	K-File COLORINOX sterile	Class IIa
A012D03100603	K-File COLORINOX sterile	Class IIa
A012D03100803	K-File COLORINOX sterile	Class IIa
A012D03101003	K-File COLORINOX sterile	Class IIa
A012D03101503	K-File COLORINOX sterile	Class IIa
A012D03102003	K-File COLORINOX sterile	Class IIa
A012D03102503	K-File COLORINOX sterile	Class IIa
A012D03103003	K-File COLORINOX sterile	Class IIa
A012D03103503	K-File COLORINOX sterile	Class IIa
A012D03104003	K-File COLORINOX sterile	Class IIa
A012D03104503	K-File COLORINOX sterile	Class IIa
A012D03105003	K-File COLORINOX sterile	Class IIa
A012D03105503	K-File COLORINOX sterile	Class IIa
A012D03106003	K-File COLORINOX sterile	Class IIa
A012D03107003	K-File COLORINOX sterile	Class IIa
A012D03108003	K-File COLORINOX sterile	Class IIa
A012D03109003	K-File COLORINOX sterile	Class IIa
A012D03110003	K-File COLORINOX sterile	Class IIa
A012D03111003	K-File COLORINOX sterile	Class IIa
A012D03112003	K-File COLORINOX sterile	Class IIa
A012D03113003	K-File COLORINOX sterile	Class IIa
A012D03114003	K-File COLORINOX sterile	Class IIa
A012D03190003	K-File COLORINOX sterile	Class IIa
A012D03190103	K-File COLORINOX sterile	Class IIa
A012D03190203	K-File COLORINOX sterile	Class IIa
A012I02101203	K-Flexofile Sterile Golden Medium	Class IIa
A012I02101703	K-Flexofile Sterile Golden Medium	Class IIa
A012I02102203	K-Flexofile Sterile Golden Medium	Class IIa
A012I02102703	K-Flexofile Sterile Golden Medium	Class IIa
A012I02103203	K-Flexofile Sterile Golden Medium	Class IIa
A012I02103703	K-Flexofile Sterile Golden Medium	Class IIa
A012I02190103	K-Flexofile Sterile Golden Medium	Class IIa
A012I02501203	K-Flexofile Sterile Golden Medium	Class IIa
A012I02501703	K-Flexofile Sterile Golden Medium	Class IIa
A012I02502203	K-Flexofile Sterile Golden Medium	Class IIa
A012I02502703	K-Flexofile Sterile Golden Medium	Class IIa
A012I02503203	K-Flexofile Sterile Golden Medium	Class IIa
A012I02503703	K-Flexofile Sterile Golden Medium	Class IIa
A012I02590103	K-Flexofile Sterile Golden Medium	Class IIa
A012I03101203	K-Flexofile Sterile Golden Medium	Class IIa
A012I03101703	K-Flexofile Sterile Golden Medium	Class IIa
A012I03102203	K-Flexofile Sterile Golden Medium	Class IIa
A012I03102703	K-Flexofile Sterile Golden Medium	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits

AOUT 2015

LNE/G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed

Item number	Product designation	Product class
A012I03103203	K-Flexofile Sterile Golden Medium	Class IIa
A012I03103703	K-Flexofile Sterile Golden Medium	Class IIa
A012I03190103	K-Flexofile Sterile Golden Medium	Class IIa
A012N02101503	K-FILE NITIFLEX STERILE	Class IIa
A012N02102003	K-FILE NITIFLEX STERILE	Class IIa
A012N02102503	K-FILE NITIFLEX STERILE	Class IIa
A012N02103003	K-FILE NITIFLEX STERILE	Class IIa
A012N02103503	K-FILE NITIFLEX STERILE	Class IIa
A012N02104003	K-FILE NITIFLEX STERILE	Class IIa
A012N02104503	K-FILE NITIFLEX STERILE	Class IIa
A012N02105003	K-FILE NITIFLEX STERILE	Class IIa
A012N02105503	K-FILE NITIFLEX STERILE	Class IIa
A012N02106003	K-FILE NITIFLEX STERILE	Class IIa
A012N02190003	K-FILE NITIFLEX STERILE	Class IIa
A012N02501503	K-FILE NITIFLEX STERILE	Class IIa
A012N02502003	K-FILE NITIFLEX STERILE	Class IIa
A012N02502503	K-FILE NITIFLEX STERILE	Class IIa
A012N02503003	K-FILE NITIFLEX STERILE	Class IIa
A012N02503503	K-FILE NITIFLEX STERILE	Class IIa
A012N02504003	K-FILE NITIFLEX STERILE	Class IIa
A012N02504503	K-FILE NITIFLEX STERILE	Class IIa
A012N02505003	K-FILE NITIFLEX STERILE	Class IIa
A012N02505503	K-FILE NITIFLEX STERILE	Class IIa
A012N02506003	K-FILE NITIFLEX STERILE	Class IIa
A012N02590003	K-FILE NITIFLEX STERILE	Class IIa
A012X01800603	C+ Files sterile for catheterization	Class IIa
A012X01800803	C+ Files sterile for catheterization	Class IIa
A012X01801003	C+ Files sterile for catheterization	Class IIa
A012X01801503	C+ Files sterile for catheterization	Class IIa
A012X01802003	C+ Files sterile for catheterization	Class IIa
A012X01890003	C+ Files sterile for catheterization	Class IIa
A012X02100603	C+ Files sterile for catheterization	Class IIa
A012X02100803	C+ Files sterile for catheterization	Class IIa
A012X02101003	C+ Files sterile for catheterization	Class IIa
A012X02101503	C+ Files sterile for catheterization	Class IIa
A012X02190003	C+ Files sterile for catheterization	Class IIa
A012X02500603	C+ Files sterile for catheterization	Class IIa
A012X02500803	C+ Files sterile for catheterization	Class IIa
A012X02501003	C+ Files sterile for catheterization	Class IIa
A012X02501503	C+ Files sterile for catheterization	Class IIa
A012X02590003	C+ Files sterile for catheterization	Class IIa
A016430000100	Endo access bur	Class IIa
A016430000200	Endo access bur	Class IIa

LNE(G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits ..... 20 AOÛT 2015

LNE(G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed

Item number	Product designation	Product class
A01643000300	Endo access bur	Class IIa
A016430090000	Endo access bur	Class IIa
A016532100100	Diamendo	Class IIa
A016532500200	Diamendo	Class IIa
A016D02100803	Hedstroem File Colorinox Sterile	Class IIa
A016D02101003	Hedstroem File Colorinox Sterile	Class IIa
A016D02101503	Hedstroem File Colorinox Sterile	Class IIa
A016D02102003	Hedstroem File Colorinox Sterile	Class IIa
A016D02102503	Hedstroem File Colorinox Sterile	Class IIa
A016D02103003	Hedstroem File Colorinox Sterile	Class IIa
A016D02103503	Hedstroem File Colorinox Sterile	Class IIa
A016D02104003	Hedstroem File Colorinox Sterile	Class IIa
A016D02104503	Hedstroem File Colorinox Sterile	Class IIa
A016D02105003	Hedstroem File Colorinox Sterile	Class IIa
A016D02105503	Hedstroem File Colorinox Sterile	Class IIa
A016D02106003	Hedstroem File Colorinox Sterile	Class IIa
A016D02107003	Hedstroem File Colorinox Sterile	Class IIa
A016D02108003	Hedstroem File Colorinox Sterile	Class IIa
A016D02109003	Hedstroem File Colorinox Sterile	Class IIa
A016D02110003	Hedstroem File Colorinox Sterile	Class IIa
A016D02111003	Hedstroem File Colorinox Sterile	Class IIa
A016D02112003	Hedstroem File Colorinox Sterile	Class IIa
A016D02113003	Hedstroem File Colorinox Sterile	Class IIa
A016D02114003	Hedstroem File Colorinox Sterile	Class IIa
A016D02190003	Hedstroem File Colorinox Sterile	Class IIa
A016D02190103	Hedstroem File Colorinox Sterile	Class IIa
A016D02190203	Hedstroem File Colorinox Sterile	Class IIa
A016D02500803	Hedstroem File Colorinox Sterile	Class IIa
A016D02501003	Hedstroem File Colorinox Sterile	Class IIa
A016D02501503	Hedstroem File Colorinox Sterile	Class IIa
A016D02502003	Hedstroem File Colorinox Sterile	Class IIa
A016D02502503	Hedstroem File Colorinox Sterile	Class IIa
A016D02503003	Hedstroem File Colorinox Sterile	Class IIa
A016D02503503	Hedstroem File Colorinox Sterile	Class IIa
A016D02504003	Hedstroem File Colorinox Sterile	Class IIa
A016D02504503	Hedstroem File Colorinox Sterile	Class IIa
A016D02505003	Hedstroem File Colorinox Sterile	Class IIa
A016D02505503	Hedstroem File Colorinox Sterile	Class IIa
A016D02506003	Hedstroem File Colorinox Sterile	Class IIa
A016D02507003	Hedstroem File Colorinox Sterile	Class IIa
A016D02508003	Hedstroem File Colorinox Sterile	Class IIa
A016D02509003	Hedstroem File Colorinox Sterile	Class IIa
A016D02510003	Hedstroem File Colorinox Sterile	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits  
 LNE/G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed

Item number	Product designation	Product class
A016D02511003	Hedstroem File Colorinox Sterile	Class IIa
A016D02512003	Hedstroem File Colorinox Sterile	Class IIa
A016D02513003	Hedstroem File Colorinox Sterile	Class IIa
A016D02514003	Hedstroem File Colorinox Sterile	Class IIa
A016D02590003	Hedstroem File Colorinox Sterile	Class IIa
A016D02590103	Hedstroem File Colorinox Sterile	Class IIa
A016D02590203	Hedstroem File Colorinox Sterile	Class IIa
A016D02800803	Hedstroem File Colorinox Sterile	Class IIa
A016D02801003	Hedstroem File Colorinox Sterile	Class IIa
A016D02801503	Hedstroem File Colorinox Sterile	Class IIa
A016D02802003	Hedstroem File Colorinox Sterile	Class IIa
A016D02802503	Hedstroem File Colorinox Sterile	Class IIa
A016D02803003	Hedstroem File Colorinox Sterile	Class IIa
A016D02803503	Hedstroem File Colorinox Sterile	Class IIa
A016D02804003	Hedstroem File Colorinox Sterile	Class IIa
A016D02804503	Hedstroem File Colorinox Sterile	Class IIa
A016D02805003	Hedstroem File Colorinox Sterile	Class IIa
A016D02805503	Hedstroem File Colorinox Sterile	Class IIa
A016D02806003	Hedstroem File Colorinox Sterile	Class IIa
A016D02807003	Hedstroem File Colorinox Sterile	Class IIa
A016D02808003	Hedstroem File Colorinox Sterile	Class IIa
A016D02809003	Hedstroem File Colorinox Sterile	Class IIa
A016D02810003	Hedstroem File Colorinox Sterile	Class IIa
A016D02811003	Hedstroem File Colorinox Sterile	Class IIa
A016D02812003	Hedstroem File Colorinox Sterile	Class IIa
A016D02813003	Hedstroem File Colorinox Sterile	Class IIa
A016D02814003	Hedstroem File Colorinox Sterile	Class IIa
A016D02890003	Hedstroem File Colorinox Sterile	Class IIa
A016D02890103	Hedstroem File Colorinox Sterile	Class IIa
A016D02890203	Hedstroem File Colorinox Sterile	Class IIa
A016D03100803	Hedstroem File Colorinox Sterile	Class IIa
A016D03101003	Hedstroem File Colorinox Sterile	Class IIa
A016D03101503	Hedstroem File Colorinox Sterile	Class IIa
A016D03102003	Hedstroem File Colorinox Sterile	Class IIa
A016D03102503	Hedstroem File Colorinox Sterile	Class IIa
A016D03103003	Hedstroem File Colorinox Sterile	Class IIa
A016D03103503	Hedstroem File Colorinox Sterile	Class IIa
A016D03104003	Hedstroem File Colorinox Sterile	Class IIa
A016D03104503	Hedstroem File Colorinox Sterile	Class IIa
A016D03105003	Hedstroem File Colorinox Sterile	Class IIa
A016D03105503	Hedstroem File Colorinox Sterile	Class IIa
A016D03106003	Hedstroem File Colorinox Sterile	Class IIa
A016D03107003	Hedstroem File Colorinox Sterile	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est  
 valide pour les dispositifs médicaux décrits  
 LNE/G-MED (0459) recognizes that its EC certificate  
 is valid for the medical devices listed

21 AOÛT 2015

Item number	Product designation	Product class
A016D03108003	Hedstroem File Colorinox Sterile	Class IIa
A016D03109003	Hedstroem File Colorinox Sterile	Class IIa
A016D03110003	Hedstroem File Colorinox Sterile	Class IIa
A016D03111003	Hedstroem File Colorinox Sterile	Class IIa
A016D03112003	Hedstroem File Colorinox Sterile	Class IIa
A016D03113003	Hedstroem File Colorinox Sterile	Class IIa
A016D03114003	Hedstroem File Colorinox Sterile	Class IIa
A016D03190003	Hedstroem File Colorinox Sterile	Class IIa
A016D03190103	Hedstroem File Colorinox Sterile	Class IIa
A016D03190203	Hedstroem File Colorinox Sterile	Class IIa
A016G02101000	File Hedstroem	Class IIa
A016G02101500	File Hedstroem	Class IIa
A016G02102000	File Hedstroem	Class IIa
A016G02102500	File Hedstroem	Class IIa
A016G02103000	File Hedstroem	Class IIa
A016G02103500	File Hedstroem	Class IIa
A016G02104000	File Hedstroem	Class IIa
A016G02104500	File Hedstroem	Class IIa
A016G02105000	File Hedstroem	Class IIa
A016G02105500	File Hedstroem	Class IIa
A016G02108000	File Hedstroem	Class IIa
A016G02190000	File Hedstroem	Class IIa
A016G02501000	File Hedstroem	Class IIa
A016G02501500	File Hedstroem	Class IIa
A016G02502000	File Hedstroem	Class IIa
A016G02502500	File Hedstroem	Class IIa
A016G02503000	File Hedstroem	Class IIa
A016G02503500	File Hedstroem	Class IIa
A016G02504000	File Hedstroem	Class IIa
A016G02504500	File Hedstroem	Class IIa
A016G02505000	File Hedstroem	Class IIa
A016G02505500	File Hedstroem	Class IIa
A016G02506000	File Hedstroem	Class IIa
A016G02507000	File Hedstroem	Class IIa
A016G02508000	File Hedstroem	Class IIa
A016G02590000	File Hedstroem	Class IIa
A016G02590100	File Hedstroem	Class IIa
A016G02801000	File Hedstroem	Class IIa
A016G02801500	File Hedstroem	Class IIa
A016G02802000	File Hedstroem	Class IIa
A016G02802500	File Hedstroem	Class IIa
A016G02803500	File Hedstroem	Class IIa
A016G02804000	File Hedstroem	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits

20 April 2015

LNE/G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed

Item number	Product designation	Product class
A017502502000	Verifier	Class IIa
A017502502003	Verifier sterile	Class IIa
A017502502500	Verifier	Class IIa
A017502502503	Verifier sterile	Class IIa
A017502503000	Verifier	Class IIa
A017502503003	Verifier sterile	Class IIa
A017502503500	Verifier	Class IIa
A017502503503	Verifier sterile	Class IIa
A017502504000	Verifier	Class IIa
A017502504003	Verifier sterile	Class IIa
A017502504500	Verifier	Class IIa
A017502504503	Verifier sterile	Class IIa
A017502505000	Verifier	Class IIa
A017502505003	Verifier sterile	Class IIa
A017502505500	Verifier	Class IIa
A017502505503	Verifier sterile	Class IIa
A017502506000	Verifier	Class IIa
A017502506003	Verifier sterile	Class IIa
A017502507000	Verifier	Class IIa
A017502507003	Verifier sterile	Class IIa
A017502508000	Verifier	Class IIa
A017502508003	Verifier sterile	Class IIa
A017502509000	Verifier	Class IIa
A017502509003	Verifier sterile	Class IIa
A017502590000	Verifier	Class IIa
A017502590003	Verifier sterile	Class IIa
A017502590100	Verifier	Class IIa
A017502590103	Verifier sterile	Class IIa
A018202100103	TAPERED FINGER SPREADER STERILE	Class IIa
A018202100203	TAPERED FINGER SPREADER STERILE	Class IIa
A018202100303	TAPERED FINGER SPREADER STERILE	Class IIa
A018202100403	TAPERED FINGER SPREADER STERILE	Class IIa
A018202190003	TAPERED FINGER SPREADER STERILE	Class IIa
A018202500103	TAPERED FINGER SPREADER STERILE	Class IIa
A018202500203	TAPERED FINGER SPREADER STERILE	Class IIa
A018202500303	TAPERED FINGER SPREADER STERILE	Class IIa
A018202500403	TAPERED FINGER SPREADER STERILE	Class IIa
A018202590003	TAPERED FINGER SPREADER STERILE	Class IIa
A022S21500300	Lentulo short, long and X-long	Class IIa
A022S21500303	Lentulo sterile Paste Carrier	Class IIa
A022S21700100	Lentulo short, long and X-long	Class IIa
A022S21700101	Lentulo sterile Paste Carrier	Class IIa
A022S21700103	Lentulo sterile Paste Carrier	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est  
 valide pour les dispositifs médicaux décrits  
 29 AOÛT 2015  
 LNE/G-MED (0459) recognizes that its EC certificate  
 is valid for the medical devices listed

Item number	Product designation	Product class
A022S21700200	Lentulo short, long and X-long	Class IIa
A022S21700203	Lentulo sterile Paste Carrier	Class IIa
A022S21700300	Lentulo short, long and X-long	Class IIa
A022S21700301	Lentulo sterile Paste Carrier	Class IIa
A022S21700303	Lentulo sterile Paste Carrier	Class IIa
A022S21700400	Lentulo short, long and X-long	Class IIa
A022S21700401	Lentulo sterile Paste Carrier	Class IIa
A022S21700403	Lentulo sterile Paste Carrier	Class IIa
A022S22100100	Lentulo short, long and X-long	Class IIa
A022S22100101	Lentulo sterile Paste Carrier	Class IIa
A022S22100103	Lentulo sterile Paste Carrier	Class IIa
A022S22100200	Lentulo short, long and X-long	Class IIa
A022S22100201	Lentulo sterile Paste Carrier	Class IIa
A022S22100203	Lentulo sterile Paste Carrier	Class IIa
A022S22100300	Lentulo short, long and X-long	Class IIa
A022S22100301	Lentulo sterile Paste Carrier	Class IIa
A022S22100303	Lentulo sterile Paste Carrier	Class IIa
A022S22100400	Lentulo short, long and X-long	Class IIa
A022S22100401	Lentulo sterile Paste Carrier	Class IIa
A022S22100403	Lentulo sterile Paste Carrier	Class IIa
A022S22500100	Lentulo short, long and X-long	Class IIa
A022S22500101	Lentulo sterile Paste Carrier	Class IIa
A022S22500103	Lentulo sterile Paste Carrier	Class IIa
A022S22500200	Lentulo short, long and X-long	Class IIa
A022S22500201	Lentulo sterile Paste Carrier	Class IIa
A022S22500203	Lentulo sterile Paste Carrier	Class IIa
A022S22500300	Lentulo short, long and X-long	Class IIa
A022S22500301	Lentulo sterile Paste Carrier	Class IIa
A022S22500303	Lentulo sterile Paste Carrier	Class IIa
A022S22500400	Lentulo short, long and X-long	Class IIa
A022S22500401	Lentulo sterile Paste Carrier	Class IIa
A022S22500403	Lentulo sterile Paste Carrier	Class IIa
A024202102500	GUTTA-CONDENSOR	Class IIa
A024202102503	GUTTA-CONDENSOR STERILE	Class IIa
A024202103000	GUTTA-CONDENSOR	Class IIa
A024202103003	GUTTA-CONDENSOR STERILE	Class IIa
A024202103500	GUTTA-CONDENSOR	Class IIa
A024202103503	GUTTA-CONDENSOR STERILE	Class IIa
A024202104000	GUTTA-CONDENSOR	Class IIa
A024202104003	GUTTA-CONDENSOR STERILE	Class IIa
A024202104500	GUTTA-CONDENSOR	Class IIa
A024202104503	GUTTA-CONDENSOR STERILE	Class IIa
A024202105000	GUTTA-CONDENSOR	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est  
 valide pour les dispositifs médicaux décrits  
 ..... 20-AOÛT 2015 .....  
 LNE/G-MED (0459) recognizes that its EC certificate  
 is valid for the medical devices listed

Item number	Product designation	Product class
A024202105003	GUTTA-CONDENSOR STERILE	Class IIa
A024202105500	GUTTA-CONDENSOR	Class IIa
A024202105503	GUTTA-CONDENSOR STERILE	Class IIa
A024202106000	GUTTA-CONDENSOR	Class IIa
A024202106003	GUTTA-CONDENSOR STERILE	Class IIa
A024202107000	GUTTA-CONDENSOR	Class IIa
A024202107003	GUTTA-CONDENSOR STERILE	Class IIa
A024202108000	GUTTA-CONDENSOR	Class IIa
A024202108003	GUTTA-CONDENSOR STERILE	Class IIa
A024202190000	GUTTA-CONDENSOR	Class IIa
A024202502500	GUTTA-CONDENSOR	Class IIa
A024202502503	GUTTA-CONDENSOR STERILE	Class IIa
A024202503000	GUTTA-CONDENSOR	Class IIa
A024202503003	GUTTA-CONDENSOR STERILE	Class IIa
A024202503500	GUTTA-CONDENSOR	Class IIa
A024202503503	GUTTA-CONDENSOR STERILE	Class IIa
A024202504000	GUTTA-CONDENSOR	Class IIa
A024202504003	GUTTA-CONDENSOR STERILE	Class IIa
A024202504500	GUTTA-CONDENSOR	Class IIa
A024202504503	GUTTA-CONDENSOR STERILE	Class IIa
A024202505000	GUTTA-CONDENSOR	Class IIa
A024202505003	GUTTA-CONDENSOR STERILE	Class IIa
A024202505500	GUTTA-CONDENSOR	Class IIa
A024202505503	GUTTA-CONDENSOR STERILE	Class IIa
A024202506000	GUTTA-CONDENSOR	Class IIa
A024202506003	GUTTA-CONDENSOR STERILE	Class IIa
A024202507000	GUTTA-CONDENSOR	Class IIa
A024202507003	GUTTA-CONDENSOR STERILE	Class IIa
A024202508000	GUTTA-CONDENSOR	Class IIa
A024202508003	GUTTA-CONDENSOR STERILE	Class IIa
A024202590000	GUTTA-CONDENSOR	Class IIa
A024402190000	GUTTA-CONDENSOR Set	Class IIa
A024402590000	GUTTA-CONDENSOR Set	Class IIa
A027400000000	Flexokit Full	Class IIa
A030500000000	Cavity Access Set	Class IIa
A030500000100	Cavity Access Set	Class IIa
A034422101500	ProFile .02	Class IIa
A034422101503	ProFile STERILE .02	Class IIa
A034422102000	ProFile .02	Class IIa
A034422102003	ProFile STERILE .02	Class IIa
A034422102500	ProFile .02	Class IIa
A034422102503	ProFile STERILE .02	Class IIa
A034422103000	ProFile .02	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits  
 2.0. ANUT 2015  
 LNE/G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed

Item number	Product designation	Product class
A034422103003	ProFile STERILE .02	Class IIa
A034422103500	ProFile .02	Class IIa
A034422103503	ProFile STERILE .02	Class IIa
A034422104000	ProFile .02	Class IIa
A034422104003	ProFile STERILE .02	Class IIa
A034422190000	ProFile .02	Class IIa
A034422190003	ProFile STERILE .02	Class IIa
A034422501500	ProFile .02	Class IIa
A034422501503	ProFile STERILE .02	Class IIa
A034422502000	ProFile .02	Class IIa
A034422502003	ProFile STERILE .02	Class IIa
A034422502500	ProFile .02	Class IIa
A034422502503	ProFile STERILE .02	Class IIa
A034422503000	ProFile .02	Class IIa
A034422503003	ProFile STERILE .02	Class IIa
A034422503500	ProFile .02	Class IIa
A034422503503	ProFile STERILE .02	Class IIa
A034422504000	ProFile .02	Class IIa
A034422504003	ProFile STERILE .02	Class IIa
A034422590000	ProFile .02	Class IIa
A034422590003	ProFile STERILE .02	Class IIa
A034521801500	ProFile .06	Class IIa
A034521801503	ProFile STERILE .06	Class IIa
A034521802000	ProFile .06	Class IIa
A034521802003	ProFile STERILE .06	Class IIa
A034521802500	ProFile .06	Class IIa
A034521802503	ProFile STERILE .06	Class IIa
A034521803000	ProFile .06	Class IIa
A034521803003	ProFile STERILE .06	Class IIa
A034521803500	ProFile .06	Class IIa
A034521803503	ProFile STERILE .06	Class IIa
A034521804000	ProFile .06	Class IIa
A034521804003	ProFile STERILE .06	Class IIa
A034521890000	ProFile .06	Class IIa
A034521890003	ProFile STERILE .06	Class IIa
A034522101500	ProFile .06	Class IIa
A034522101503	ProFile STERILE .06	Class IIa
A034522102000	ProFile .06	Class IIa
A034522102003	ProFile STERILE .06	Class IIa
A034522102500	ProFile .06	Class IIa
A034522102503	ProFile STERILE .06	Class IIa
A034522103000	ProFile .06	Class IIa
A034522103003	ProFile STERILE .06	Class IIa

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 valide pour les dispositifs médicaux décrits  
 LNE/G-MED (0459) recognizes that its EC certificate  
 is valid for the medical devices listed  
 20...AOUT 2015

Item number	Product designation	Product class
A034522103500	ProFile .06	Class IIa
A034522103503	ProFile STERILE .06	Class IIa
A034522104000	ProFile .06	Class IIa
A034522104003	ProFile STERILE .06	Class IIa
A034522190000	ProFile .06	Class IIa
A034522190003	ProFile STERILE .06	Class IIa
A034522501500	ProFile .06	Class IIa
A034522501503	ProFile STERILE .06	Class IIa
A034522502000	ProFile .06	Class IIa
A034522502003	ProFile STERILE .06	Class IIa
A034522502500	ProFile .06	Class IIa
A034522502503	ProFile STERILE .06	Class IIa
A034522503000	ProFile .06	Class IIa
A034522503003	ProFile STERILE .06	Class IIa
A034522503500	ProFile .06	Class IIa
A034522503503	ProFile STERILE .06	Class IIa
A034522504000	ProFile .06	Class IIa
A034522504003	ProFile STERILE .06	Class IIa
A034522590000	ProFile .06	Class IIa
A034522590003	ProFile STERILE .06	Class IIa
A035802590000	ProFile O.S. + .06 + .04	Class IIa
A035921900100	ProFile O.S.	Class IIa
A035921900103	ProFile STERILE O.S.	Class IIa
A035921900200	ProFile O.S.	Class IIa
A035921900203	ProFile STERILE O.S.	Class IIa
A035921900300	ProFile O.S.	Class IIa
A035921900303	ProFile STERILE O.S.	Class IIa
A035921900400	ProFile O.S.	Class IIa
A035921900403	ProFile STERILE O.S.	Class IIa
A035921900500	ProFile O.S.	Class IIa
A035921900503	ProFile STERILE O.S.	Class IIa
A035921900600	ProFile O.S.	Class IIa
A035921900603	ProFile STERILE O.S.	Class IIa
A035921990000	ProFile O.S.	Class IIa
A035921990003	ProFile STERILE O.S.	Class IIa
A036102101000	Unifile	Class IIa
A036102101500	Unifile LNE/G-MED (0459) reconnaît que son certificat CE est	Class IIa
A036102102000	Unifile valide pour les dispositifs médicaux décrits	Class IIa
A036102102500	Unifile ..... 2015	Class IIa
A036102103000	Unifile LNE/G-MED (0459) recognizes that its EC certificate	Class IIa
A036102103500	Unifile is valid for the medical devices listed	Class IIa
A036102105000	Unifile	Class IIa
A036102105500	Unifile	Class IIa

Item number	Product designation	Product class
A036102106000	Unifile	Class IIa
A036102107000	Unifile	Class IIa
A036102190000	Unifile	Class IIa
A036102190100	Unifile	Class IIa
A036102191000	Unifile	Class IIa
A036102501000	Unifile	Class IIa
A036102501500	Unifile	Class IIa
A036102502000	Unifile	Class IIa
A036102503000	Unifile	Class IIa
A036102504000	Unifile	Class IIa
A036102504500	Unifile	Class IIa
A036102505000	Unifile	Class IIa
A036102505500	Unifile	Class IIa
A036102506000	Unifile	Class IIa
A036102507000	Unifile	Class IIa
A036102508000	Unifile	Class IIa
A036102590000	Unifile	Class IIa
A036102590100	Unifile	Class IIa
A036102591000	Unifile	Class IIa
A036200000000	PROFILE Organizer	Class IIa
A036200000900	PROFILE Organizer	Class IIa
A036302100100	PROFILE	Class IIa
A036302500200	PROFILE	Class IIa
A040922100100	PROTAPER UNIVERSAL SEQ BASE	Class IIa
A040922100103	PROTAPER UNIVERSAL Sterile assortment SEQ BASE	Class IIa
A040922190100	PROTAPER UNIVERSAL SEQ BASE	Class IIa
A040922190103	PROTAPER UNIVERSAL Sterile assortment SEQ BASE	Class IIa
A040922500200	PROTAPER UNIVERSAL SEQ BASE	Class IIa
A040922500203	PROTAPER UNIVERSAL Sterile assortment SEQ BASE	Class IIa
A040922590100	PROTAPER UNIVERSAL SEQ BASE	Class IIa
A040922590103	PROTAPER UNIVERSAL Sterile assortment SEQ BASE	Class IIa
A040923100300	PROTAPER UNIVERSAL SEQ BASE	Class IIa
A040923100303	PROTAPER UNIVERSAL Sterile assortment SEQ BASE	Class IIa
A040923190100	PROTAPER UNIVERSAL SEQ BASE	Class IIa
A040923190103	PROTAPER UNIVERSAL Sterile assortment SEQ BASE	Class IIa
A041021910100	PROTAPER UNIVERSAL Shaping Files	Class IIa
A041021910103	PROTAPER UNIVERSAL Shaping Files Sterile	Class IIa
A041022110100	PROTAPER UNIVERSAL Shaping Files	Class IIa
A041022110103	PROTAPER UNIVERSAL Shaping Files Sterile	Class IIa
A041022110200	PROTAPER UNIVERSAL Shaping Files	Class IIa
A041022110203	PROTAPER UNIVERSAL Shaping Files Sterile	Class IIa
A041022510100	PROTAPER UNIVERSAL Shaping Files	Class IIa
A041022510103	PROTAPER UNIVERSAL Shaping Files Sterile	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits  
 LNE/G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed

Item number	Product designation	Product class
A041022510200	PROTAPER UNIVERSAL Shaping Files	Class IIa
A041022510203	PROTAPER UNIVERSAL Shaping Files Sterile	Class IIa
A041023110100	PROTAPER UNIVERSAL Shaping Files	Class IIa
A041023110103	PROTAPER UNIVERSAL Shaping Files Sterile	Class IIa
A041023110200	PROTAPER UNIVERSAL Shaping Files	Class IIa
A041023110203	PROTAPER UNIVERSAL Shaping Files Sterile	Class IIa
A041122110100	PROTAPER UNIVERSAL Finishing Files	Class IIa
A041122110103	PROTAPER UNIVERSAL Finishing Files Sterile	Class IIa
A041122110200	PROTAPER UNIVERSAL Finishing Files	Class IIa
A041122110203	PROTAPER UNIVERSAL Finishing Files Sterile	Class IIa
A041122110300	PROTAPER UNIVERSAL Finishing Files	Class IIa
A041122110303	PROTAPER UNIVERSAL Finishing Files Sterile	Class IIa
A041122110400	PROTAPER UNIVERSAL Finishing Files	Class IIa
A041122110403	PROTAPER UNIVERSAL Finishing Files Sterile	Class IIa
A041122110500	PROTAPER UNIVERSAL Finishing Files	Class IIa
A041122110503	PROTAPER UNIVERSAL Finishing Files Sterile	Class IIa
A041122510100	PROTAPER UNIVERSAL Finishing Files	Class IIa
A041122510103	PROTAPER UNIVERSAL Finishing Files Sterile	Class IIa
A041122510200	PROTAPER UNIVERSAL Finishing Files	Class IIa
A041122510203	PROTAPER UNIVERSAL Finishing Files Sterile	Class IIa
A041122510300	PROTAPER UNIVERSAL Finishing Files	Class IIa
A041122510303	PROTAPER UNIVERSAL Finishing Files Sterile	Class IIa
A041122510400	PROTAPER UNIVERSAL Finishing Files	Class IIa
A041122510403	PROTAPER UNIVERSAL Finishing Files Sterile	Class IIa
A041122510500	PROTAPER UNIVERSAL Finishing Files	Class IIa
A041122510503	PROTAPER UNIVERSAL Finishing Files Sterile	Class IIa
A041123110100	PROTAPER UNIVERSAL Finishing Files	Class IIa
A041123110103	PROTAPER UNIVERSAL Finishing Files Sterile	Class IIa
A041123110200	PROTAPER UNIVERSAL Finishing Files	Class IIa
A041123110203	PROTAPER UNIVERSAL Finishing Files Sterile	Class IIa
A041123110300	PROTAPER UNIVERSAL Finishing Files	Class IIa
A041123110303	PROTAPER UNIVERSAL Finishing Files Sterile	Class IIa
A041123110400	PROTAPER UNIVERSAL Finishing Files	Class IIa
A041123110403	PROTAPER UNIVERSAL Finishing Files Sterile	Class IIa
A041123110500	PROTAPER UNIVERSAL Finishing Files	Class IIa
A041123110503	PROTAPER UNIVERSAL Finishing Files Sterile	Class IIa
A041522190100	Starter Kit PROTAPER UNIVERSAL	Class IIa
A041522590100	Starter Kit PROTAPER UNIVERSAL	Class IIa
A041523190100	Starter Kit PROTAPER UNIVERSAL	Class IIa
A066000000000	START-X®	Class IIa
A066000000100	START-X®	Class IIa
A066000000200	START-X® LNE/G-MED (0459) reconnaît que son certificat CE est	Class IIa
A066000000300	START-X® valide pour les dispositifs médicaux décrits	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits

Item number	Product designation	Product class
A066000000400	START-X®	Class IIa
A066000000500	START-X®	Class IIa
A066000090000	START-X®	Class IIa
A066100000100	START-X®	Class IIa
A066100000200	START-X® LNE/G-MED (0459) reconnaît que son certificat CE est	Class IIa
A066100000300	START-X® valide pour les dispositifs médicaux décrits	Class IIa
A066100000400	START-X® LNE/G-MED (0459) recognizes that its EC certificate	Class IIa
A066100000500	START-X® is valid for the medical devices listed	Class IIa
A066100090000	START-X®	Class IIa
A070022100L01	WAVEONE reciprocating file sterile	Class IIa
A070022100P01	WAVEONE reciprocating file sterile	Class IIa
A070022100S01	WAVEONE reciprocating file sterile	Class IIa
A070022190A01	WAVEONE Assortment Reciprocating Files Sterile	Class IIa
A070022500L01	WAVEONE reciprocating file sterile	Class IIa
A070022500P01	WAVEONE reciprocating file sterile	Class IIa
A070022500S01	WAVEONE reciprocating file sterile	Class IIa
A070022590A01	WAVEONE Assortment Reciprocating Files Sterile	Class IIa
A070023100L01	WAVEONE reciprocating file sterile	Class IIa
A070023100P01	WAVEONE reciprocating file sterile	Class IIa
A070023100S01	WAVEONE reciprocating file sterile	Class IIa
A070023190A01	WAVEONE Assortment Reciprocating Files Sterile	Class IIa
A070622100L01	WAVEONE reciprocating file sterile	Class IIa
A070622100P01	WAVEONE reciprocating file sterile	Class IIa
A070622100S01	WAVEONE reciprocating file sterile	Class IIa
A070622190A01	WAVEONE Assortment Reciprocating Files Sterile	Class IIa
A070622500L01	WAVEONE reciprocating file sterile	Class IIa
A070622500P01	WAVEONE reciprocating file sterile	Class IIa
A070622500S01	WAVEONE reciprocating file sterile	Class IIa
A070622590A01	WAVEONE Assortment Reciprocating Files Sterile	Class IIa
A070623100L01	WAVEONE reciprocating file sterile	Class IIa
A070623100P01	WAVEONE reciprocating file sterile	Class IIa
A070623100S01	WAVEONE reciprocating file sterile	Class IIa
A070623190A01	WAVEONE Assortment Reciprocating Files Sterile	Class IIa
A0751225G9D03	WAVEONE GOLD One Reciprocating Files Sterile	Class IIa
A0753221G0L03	WAVEONE GOLD Reciprocating Files Sterile	Class IIa
A0753221G0M03	WAVEONE GOLD Reciprocating Files Sterile	Class IIa
A0753221G0P03	WAVEONE GOLD Reciprocating Files Sterile	Class IIa
A0753221G0S03	WAVEONE GOLD Reciprocating Files Sterile	Class IIa
A0753225G0L03	WAVEONE GOLD Reciprocating Files Sterile	Class IIa
A0753225G0M03	WAVEONE GOLD Reciprocating Files Sterile	Class IIa
A0753225G0P03	WAVEONE GOLD Reciprocating Files Sterile	Class IIa
A0753225G0S03	WAVEONE GOLD Reciprocating Files Sterile	Class IIa
A0753231G0L03	WAVEONE GOLD Reciprocating Files Sterile	Class IIa

Item number	Product designation	Product class
A0753231G0M03	WAVEONE GOLD Reciprocating Files Sterile	Class IIa
A0753231G0P03	WAVEONE GOLD Reciprocating Files Sterile	Class IIa
A0753231G0S03	WAVEONE GOLD Reciprocating Files Sterile	Class IIa
A0754221G9A03	WAVEONE GOLD Assortment Reciprocating Files Sterile	Class IIa
A0754225G9A03	WAVEONE GOLD Assortment Reciprocating Files Sterile	Class IIa
A0754231G9A03	WAVEONE GOLD Assortment Reciprocating Files Sterile	Class IIa
A0756221G0L03	WAVEONE GOLD Reciprocating Files Sterile	Class IIa
A0756221G0M03	WAVEONE GOLD Reciprocating Files Sterile	Class IIa
A0756221G0P03	WAVEONE GOLD Reciprocating Files Sterile	Class IIa
A0756221G0S03	WAVEONE GOLD Reciprocating Files Sterile	Class IIa
A0756225G0L03	WAVEONE GOLD Reciprocating Files Sterile	Class IIa
A0756225G0M03	WAVEONE GOLD Reciprocating Files Sterile	Class IIa
A0756225G0P03	WAVEONE GOLD Reciprocating Files Sterile	Class IIa
A0756225G0S03	WAVEONE GOLD Reciprocating Files Sterile	Class IIa
A0756231G0L03	WAVEONE GOLD Reciprocating Files Sterile	Class IIa
A0756231G0M03	WAVEONE GOLD Reciprocating Files Sterile	Class IIa
A0756231G0P03	WAVEONE GOLD Reciprocating Files Sterile	Class IIa
A0756231G0S03	WAVEONE GOLD Reciprocating Files Sterile	Class IIa
A080021900A03	PROTAPER NEXT XA	Class IIa
A080022100103	PROTAPER NEXT	Class IIa
A080022100203	PROTAPER NEXT	Class IIa
A080022100303	PROTAPER NEXT	Class IIa
A080022100403	PROTAPER NEXT	Class IIa
A080022100503	PROTAPER NEXT	Class IIa
A08002219A003	PROTAPER NEXT Assortment	Class IIa
A080022500103	PROTAPER NEXT	Class IIa
A080022500203	PROTAPER NEXT	Class IIa
A080022500303	PROTAPER NEXT	Class IIa
A080022500403	PROTAPER NEXT	Class IIa
A080022500503	PROTAPER NEXT	Class IIa
A08002259A003	PROTAPER NEXT Assortment	Class IIa
A080023100103	PROTAPER NEXT	Class IIa
A080023100203	PROTAPER NEXT	Class IIa
A080023100303	PROTAPER NEXT	Class IIa
A080023100403	PROTAPER NEXT	Class IIa
A080023100503	PROTAPER NEXT	Class IIa
A08002319A003	PROTAPER NEXT Assortment	Class IIa
A08022259D003	PROTAPER NEXT Demo	Class IIa
A080321900A03	PROTAPER NEXT XA	Class IIa
A080322100103	PROTAPER NEXT	Class IIa
A080322100203	PROTAPER NEXT	Class IIa
A080322100303	PROTAPER NEXT	Class IIa
A080322100403	PROTAPER NEXT	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est  
 valide pour les dispositifs médicaux décrits  
 .....  
 LNE/G-MED (0459) recognizes that its EC certificate  
 is valid for the medical devices listed

Item number	Product designation	Product class
A080322100503	PROTAPER NEXT	Class IIa
A08032219A003	PROTAPER NEXT Assortment	Class IIa
A080322500103	PROTAPER NEXT	Class IIa
A080322500203	PROTAPER NEXT	Class IIa
A080322500303	PROTAPER NEXT	Class IIa
A080322500403	PROTAPER NEXT	Class IIa
A080322500503	PROTAPER NEXT	Class IIa
A08032259A003	PROTAPER NEXT Assortment	Class IIa
A080323100103	PROTAPER NEXT	Class IIa
A080323100203	PROTAPER NEXT	Class IIa
A080323100303	PROTAPER NEXT	Class IIa
A080323100403	PROTAPER NEXT	Class IIa
A080323100503	PROTAPER NEXT	Class IIa
A08032319A003	PROTAPER NEXT Assortment	Class IIa
A082100023100	PRORINSE	Class IIa
A082100023400	PRORINSE	Class IIa
A082100024100	PRORINSE	Class IIa
A082100024400	PRORINSE	Class IIa
A082100025100	PRORINSE	Class IIa
A082100025400	PRORINSE	Class IIa
A082100028100	PRORINSE	Class IIa
A082100028400	PRORINSE	Class IIa
A082100030100	PRORINSE	Class IIa
A082100030400	PRORINSE	Class IIa
A092322100103	PROGLIDER is valid for the medical devices listed	Class IIa
A092322500103	PROGLIDER	Class IIa
A092323100103	PROGLIDER	Class IIa
A09242259D003	PROGLIDER	Class IIa
A092622100103	PROGLIDER	Class IIa
A092622500103	PROGLIDER	Class IIa
A092623100103	PROGLIDER	Class IIa
A100400000000	X-SMART	Class IIa
A100400000022	X-SMART	Class IIa
A100400000033	X-SMART	Class IIa
A100400000122	X-SMART	Class IIa
A100400000126	X-SMART	Class IIa
A100400000133	X-SMART	Class IIa
A101800000000	WAVEONE endo motor	Class IIa
A101800000200	WAVEONE endo motor	Class IIa
A101800000400	WAVEONE endo motor	Class IIa
A102400000033	X-SMART DUAL	Class IIa
A102800000026	PROPEX II	Class IIa
A102800000033	PROPEX II	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits  
 2 D'AOÛT 2015  
 LNE/G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed

Item number	Product designation	Product class
A10300000000	PROPEX PIXI™	Class IIa
A10300000100	PROPEX PIXI™	Class IIa
A141021600100	ProTaper Universal Retreatment	Class IIa
A141021600103	ProTaper Universal Retreatment Sterile	Class IIa
A141021800100	ProTaper Universal Retreatment	Class IIa
A141021800103	ProTaper Universal Retreatment Sterile	Class IIa
A141022200100	ProTaper Universal Retreatment	Class IIa
A141022200103	ProTaper Universal Retreatment Sterile	Class IIa
A141220090000	ProTaper Universal Retreatment	Class IIa
A141220090003	ProTaper Universal Retreatment	Class IIa
A141500090000	ProTaper Universal Starter Kit	Class IIa
31050701A0000	X-SMART IQ HANDPIECE MOTOR	Class IIa
A105100000000	X-SMART IQ CONTRA-ANGLE	Class IIa
A230A00000200	Endo Set Univ. Fernando Pessoa	Class IIa
A230A00000300	Endoset Universidad catolica portugeses polo de viseu	Class IIa
A230A00000600	Endoset Universidad del Pais vasco	Class IIa
A230A00000700	Endoset Universidad de Zaragoza	Class IIa
A230A00001400	Endoset ISCS-EGAS Moniz	Class IIa
A230A00001500	Endoset Universidad San Pablo CEU de Madrid	Class IIa
A230A00002000	ENDO SET UNIVERSIDAD EUROPE MIGUEL DE CERVANTES	Class IIa
A305Z00000000	Cavity Access Z Set	Class IIa
A305Z00000100	Cavity Access Z Set	Class IIa
A966000090100	START-X® Trial Kit Compatible Satelec	Class IIa
A966100090100	START-X® Trial Kit Compatible EMS	Class IIa
C005320000100	MOOSER SYSTEM	Class IIa
C005320000200	MOOSER SYSTEM	Class IIa
C005420000100	MOOSER SYSTEM	Class IIa
C005420000200	MOOSER SYSTEM	Class IIa
C005820000000	STP RESTORATIVE PINS	Class IIa
C009800000100	RS-RADIX-STIFTE INTEGRAL SET	Class IIa
C009800000200	RS-RADIX-STIFTE INTEGRAL SET	Class IIa
C009800000300	RS-RADIX-STIFTE INTEGRAL SET	Class IIa
C009900000000	RS-RADIX-STIFTE SET	Class IIa
C010000000100	RS RADIX	Class IIa
C010000000200	RS RADIX	Class IIa
C010000000300	RS RADIX	Class IIa
C010120000100	RS RADIX LNE/G-MED (0459) reconnaît que son certificat CE est	Class IIa
C010120000200	RS RADIX valide pour les dispositifs médicaux décrits	Class IIa
C010120000300	RS RADIX LNE/G-MED (0459) recognizes that its EC certificate	Class IIa
C010220000100	RS RADIX is valid for the medical devices listed	Class IIa
C010220000200	RS RADIX	Class IIa
C010220000300	RS RADIX	Class IIa
C019300000100	RADIX-ANKER-STANDARD	Class IIa

Item number	Product designation	Product class
C01930000200	RADIX-ANKER-STANDARD	Class IIa
C01930000300	RADIX-ANKER-STANDARD	Class IIa
C02000000000	STP RESTORATIVE PINS	Class IIa
C02020080000	MOOSER SYSTEM INTEGRAL SET	Class IIa
C020200800100	MOOSER SYSTEM INTEGRAL SET	Class IIa
C020201000000	MOOSER SYSTEM INTEGRAL SET	Class IIa
C020201000100	MOOSER SYSTEM INTEGRAL SET	Class IIa
C020820000L00	STP restorative pins	Class IIa
C020820000S00	STP restorative pins	Class IIa
C02100000000	STP RESTORATIVE PINS INTEGRAL SET	Class IIa
C02110000000	STP-REFILL	Class IIa
C021220000800	UNIMETRIC X mm	Class IIa
C021220010800	UNIMETRIC X mm	Class IIa
C021220011000	UNIMETRIC X mm	Class IIa
C021220020800	UNIMETRIC X mm	Class IIa
C021220021000	UNIMETRIC X mm	Class IIa
C021220030800	UNIMETRIC X mm	Class IIa
C021220031000	UNIMETRIC X mm	Class IIa
C021320000800	UNIMETRIC X mm	Class IIa
C021320010800	UNIMETRIC X mm	Class IIa
C021320011000	UNIMETRIC X mm	Class IIa
C021320020800	UNIMETRIC X mm	Class IIa
C021320021000	UNIMETRIC X mm	Class IIa
C021320030800	UNIMETRIC X mm	Class IIa
C021320031000	UNIMETRIC X mm	Class IIa
C021420000000	UNIMETRIC	Class IIa
C021620000100	RADIX-ANKER-STANDARD	Class IIa
C021620000200	RADIX-ANKER-STANDARD	Class IIa
C021620000300	RADIX-ANKER-STANDARD	Class IIa
C0216200003B00	RADIX-ANKER-STANDARD	Class IIa
C021720000100	RADIX-ANKER-STANDARD	Class IIa
C021720000200	RADIX-ANKER-STANDARD	Class IIa
C021720000300	RADIX-ANKER-STANDARD	Class IIa
C0217200003B00	RADIX-ANKER-STANDARD	Class IIa
C022020000100	RADIX-ANKER-STANDARD	Class IIa
C022020000200	RADIX-ANKER-STANDARD	Class IIa
C022020000300	RADIX-ANKER-STANDARD	Class IIa
C0220200003B00	RADIX-ANKER-STANDARD	Class IIa
C022500000000	STP RESTORATIVE PIN	Class IIa
C024100000800	Mooser System	Class IIa
C024100001000	Mooser System	Class IIa
C024300000000	Mooser system Integral set	Class IIa
C026100000100	Radix-Anker-Long	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits

LNE/G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed

2015

Item number	Product designation	Product class
C02610000200	Radix-Anker-Long	Class IIa
C02610000300	Radix-Anker-Long	Class IIa
C02632000100	Radix-Anker-Long	Class IIa
C02632000200	Radix-Anker-Long	Class IIa
C02632000300	Radix-Anker-Long	Class IIa
C031000000000	Radix-Anker-Long set	Class IIa
C03200000200	Radix-Anker-Standard Demo kit	Class IIa
C032100020800	Unimetric T demo kit	Class IIa
C032100021000	Unimetric T demo kit	Class IIa
C03220000200	Radix-Anker-Long demo kit	Class IIa
C032420000000	Radix Anker Long	Class IIa
C055B00000100	MOOSER SYSTEM	Class IIa
C055B00000200	MOOSER SYSTEM	Class IIa
C055B00000300	MOOSER SYSTEM	Class IIa
C055B00000400	MOOSER SYSTEM	Class IIa
C055C00950800	MOOSER SYSTEM	Class IIa
C055C00951000	MOOSER SYSTEM	Class IIa
C055C01150800	MOOSER SYSTEM	Class IIa
C055C01151000	MOOSER SYSTEM	Class IIa
C055C01350800	MOOSER SYSTEM	Class IIa
C055C01351000	MOOSER SYSTEM	Class IIa
C055D00000800	MOOSER SYSTEM	Class IIa
C055D00001000	MOOSER SYSTEM	Class IIa
C055D01200800	MOOSER SYSTEM	Class IIa
C055D01400800	MOOSER SYSTEM	Class IIa
C055D01401000	MOOSER SYSTEM	Class IIa
C055D01601000	MOOSER SYSTEM	Class IIa
C060000000100	EasyPost	Class IIa
C060000000200	EasyPost	Class IIa
C060000000300	EasyPost	Class IIa
C060000000400	EasyPost	Class IIa
C060000000500	EasyPost	Class IIa
C060000000600	EasyPost	Class IIa
C060100000100	Easy Post	Class IIa
C060100000200	Easy Post	Class IIa
C060100000300	Easy Post	Class IIa
C060100000400	Easy Post	Class IIa
C060100000500	Easy Post	Class IIa
C060100000600	Easy Post	Class IIa
C060200800100	EasyPost	Class IIa
C060201000100	EasyPost	Class IIa
C060201300100	EasyPost	Class IIa
C061300000100	Radix Fiber Post	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits  
 20 AOÛT 2015  
 LNE/G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed

Item number	Product designation	Product class
C06130000200	Radix Fiber Post	Class IIa
C06130000300	Radix Fiber Post	Class IIa
C06130000400	Radix Fiber Post	Class IIa
C06130000500	Radix Fiber Post	Class IIa
C06130000600	Radix Fiber Post	Class IIa
C061400800100	Radix Fiber Post Basic Kit	Class IIa
C061400800200	Radix Fiber Post Trial Kit	Class IIa
C061401000100	Radix Fiber Post Basic Kit	Class IIa
C061401000200	Radix Fiber Post Trial Kit	Class IIa
C061401300100	Radix Fiber Post Basic Kit	Class IIa
C106KL0050A00	CYTCO-K	Class IIa
C106KL0050B00	CYTCO-K	Class IIa
C106KL0080A00	CYTCO-K	Class IIa
C106KL0080B00	CYTCO-K	Class IIa
C106KS0050A00	CYTCO-K	Class IIa
C107KL0005000	CYTCO-K	Class IIa
C107KL0008000	CYTCO-K LNE/G-MED (0459) reconnaît que son certificat CE est	Class IIa
C107KS0005000	CYTCO-K valide pour les dispositifs médicaux décrits	Class IIa
C108KL0005000	CYTCO-K	Class IIa
C108KL0008000	CYTCO-K LNE/G-MED (0459) recognizes that its EC certificate	Class IIa
C108KS0005000	CYTCO-K is valid for the medical devices listed	Class IIa
C117K00000000	CYTCO-K SET	Class IIa
C117K00000100	CYTCO-K SET	Class IIa
C118KL5000100	CYTCO-K INTEGRAL SET	Class IIa
C118KL5000200	CYTCO-K INTEGRAL SET	Class IIa
C118KL8000100	CYTCO-K INTEGRAL SET	Class IIa
C118KL8000200	CYTCO-K INTEGRAL SET	Class IIa
C118KS5000100	CYTCO-K INTEGRAL SET	Class IIa
C193T00000100	RADIX-ANKER-STANDARD	Class Is
C193T00000200	RADIX-ANKER-STANDARD	Class Is
C193T00000300	RADIX-ANKER-STANDARD	Class Is
C193T000003B00	RADIX-ANKER-STANDARD	Class Is
C195C00000100	RADIX-ANKER-STANDARD INTEGRAL SET	Class IIa
C195C00000200	RADIX-ANKER-STANDARD INTEGRAL SET	Class IIa
C195C00000300	RADIX-ANKER-STANDARD INTEGRAL SET	Class IIa
C195CT0000100	RADIX-ANKER-STANDARD INTEGRAL SET	Class IIa
C195CT0000200	RADIX-ANKER-STANDARD INTEGRAL SET	Class IIa
C195CT0000300	RADIX-ANKER-STANDARD INTEGRAL SET	Class IIa
C195CT00003B00	RADIX-ANKER-STANDARD INTEGRAL SET	Class IIa
C195CTR000100	RADIX-ANKER-STANDARD INTEGRAL SET	Class IIa
C195CTR000200	RADIX-ANKER-STANDARD INTEGRAL SET	Class IIa
C195CTR000300	RADIX-ANKER-STANDARD INTEGRAL SET	Class IIa
C195D00000000	RADIX-ANKER-STANDARD SET	Class IIa

Item number	Product designation	Product class
C195DT0000000	RADIX-ANKER-STANDARD SET	Class IIa
C195ET0000000	RADIX-ANKER-STANDARD SET	Class IIa
C211R00000000	STP RESTORATIVE PINS INTEGRAL SET	Class IIa
C215TL0010800	UNIMETRIC X.X MM	Class IIa
C215TL0011000	UNIMETRIC X.X MM	Class IIa
C215TL0020800	UNIMETRIC X.X MM	Class IIa
C215TL0021000	UNIMETRIC X.X MM	Class IIa
C215TL0030800	UNIMETRIC X.X MM	Class IIa
C215TL0031000	UNIMETRIC X.X MM	Class IIa
C215TRS020800	UNIMETRIC X.X MM	Class IIa
C215TS0000800	UNIMETRIC X.X MM	Class IIa
C215TS0010800	UNIMETRIC X.X MM	Class IIa
C215TS0011000	UNIMETRIC X.X MM	Class IIa
C215TS0020800	UNIMETRIC X.X MM	Class IIa
C215TS0021000	UNIMETRIC X.X MM	Class IIa
C215TS0030800	UNIMETRIC X.X MM	Class IIa
C215TS0031000	UNIMETRIC X.X MM	Class IIa
C215U00010800	UNICLIP	Class IIa
C215U00011000	UNICLIP LNE/G-MED (0459) reconnaît que son certificat CE est	Class IIa
C215U00020800	UNICLIP valide pour les dispositifs médicaux décrits	Class IIa
C215U00021000	UNICLIP LNE/G-MED (0459) recognizes that its EC certificate	Class IIa
C215U00030800	UNICLIP is valid for the medical devices listed	Class IIa
C215U00031000	UNICLIP	Class IIa
C226T00000800	UNIMETRIC XMM SET	Class IIa
C226T00001000	UNIMETRIC XMM SET	Class IIa
C226TR0000800	UNIMETRIC XMM SET	Class IIa
C226TR0001000	UNIMETRIC XMM SET	Class IIa
C226U00000800	UNICLIP XMM SET	Class IIa
C226U00001000	UNICLIP XMM SET	Class IIa
D002314000500	Steel burs cavity round	Class IIa
D002314000600	Steel burs cavity round	Class IIa
D002314000700	Steel burs cavity round	Class IIa
D002314000800	Steel burs cavity round	Class IIa
D002314000900	Steel burs cavity round	Class IIa
D002314001000	Steel burs cavity round	Class IIa
D002314001200	Steel burs cavity round	Class IIa
D002314001400	Steel burs cavity round	Class IIa
D002314001600	Steel burs cavity round	Class IIa
D002314001800	Steel burs cavity round	Class IIa
D002314002100	Steel burs cavity round	Class IIa
D002314002300	Steel burs cavity round	Class IIa
D002314002500	Steel burs cavity round	Class IIa
D002314002700	Steel burs cavity round	Class IIa

Item number	Product designation	Product class
D002314002900	Steel burs cavity round	Class IIa
D002314003100	Steel burs cavity round	Class IIa
D002314003300	Steel burs cavity round	Class IIa
D002314003500	Steel burs cavity round	Class IIa
D002314003700	Steel burs cavity round	Class IIa
D002314004000	Steel burs cavity round	Class IIa
D002314004200	Steel burs cavity round	Class IIa
D002324000500	Steel burs cavity round	Class IIa
D002324000600	Steel burs cavity round	Class IIa
D002324000700	Steel burs cavity round	Class IIa
D002324000800	Steel burs cavity round	Class IIa
D002324000900	Steel burs cavity round	Class IIa
D002324001000	Steel burs cavity round	Class IIa
D002324001200	Steel burs cavity round	Class IIa
D002324001400	Steel burs cavity round	Class IIa
D002324001600	Steel burs cavity round	Class IIa
D002324001800	Steel burs cavity round	Class IIa
D002324002100	Steel burs cavity round	Class IIa
D002324002300	Steel burs cavity round	Class IIa
D002324002500	Steel burs cavity round	Class IIa
D002324002700	Steel burs cavity round	Class IIa
D002324002900	Steel burs cavity round	Class IIa
D002324003100	Steel burs cavity round	Class IIa
D002324003300	Steel burs cavity round	Class IIa
D002324003500	Steel burs cavity round	Class IIa
D002324003700	Steel burs cavity round	Class IIa
D002324004000	Steel burs cavity round	Class IIa
D002324004200	Steel burs cavity round	Class IIa
D002325000600	Steel burs cavity round	Class IIa
D002325000700	Steel burs cavity round	Class IIa
D002325000800	Steel burs cavity round	Class IIa
D002325000900	Steel burs cavity round	Class IIa
D002325001000	Steel burs cavity round	Class IIa
D002325001200	Steel burs cavity round	Class IIa
D002325001400	Steel burs cavity round	Class IIa
D002325001600	Steel burs cavity round	Class IIa
D002325001800	Steel burs cavity round	Class IIa
D002325002100	Steel burs cavity round	Class IIa
D002325002300	Steel burs cavity round	Class IIa
D002325002500	Steel burs cavity round	Class IIa
D002325002700	Steel burs cavity round	Class IIa
D002325002900	Steel burs cavity round	Class IIa
D002325003100	Steel burs cavity round	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est  
 valide pour les dispositifs médicaux décrits  
 20 AOÛT 2019  
 LNE/G-MED (0459) recognizes that its EC certificate  
 is valid for the medical devices listed

Item number	Product designation	Product class
D002325003300	Steel burs cavity round	Class IIa
D002325003500	Steel burs cavity round	Class IIa
D002325003700	Steel burs cavity round	Class IIa
D002325004000	Steel burs cavity round	Class IIa
D002325004200	Steel burs cavity round	Class IIa
D002414000600	Steel burs cavity inverted cone	Class IIa
D002414000700	Steel burs cavity inverted cone	Class IIa
D002414000800	Steel burs cavity inverted cone	Class IIa
D002414000900	Steel burs cavity inverted cone	Class IIa
D002414001000	Steel burs cavity inverted cone	Class IIa
D002414001200	Steel burs cavity inverted cone	Class IIa
D002414001400	Steel burs cavity inverted cone	Class IIa
D002414001600	Steel burs cavity inverted cone	Class IIa
D002414001800	Steel burs cavity inverted cone	Class IIa
D002414002100	Steel burs cavity inverted cone	Class IIa
D002414002300	Steel burs cavity inverted cone	Class IIa
D002414002500	Steel burs cavity inverted cone	Class IIa
D002414002700	Steel burs cavity inverted cone	Class IIa
D002414002900	Steel burs cavity inverted cone	Class IIa
D002424000600	Steel burs cavity inverted cone	Class IIa
D002424000700	Steel burs cavity inverted cone	Class IIa
D002424000800	Steel burs cavity inverted cone	Class IIa
D002424000900	Steel burs cavity inverted cone	Class IIa
D002424001000	Steel burs cavity inverted cone	Class IIa
D002424001200	Steel burs cavity inverted cone	Class IIa
D002424001400	Steel burs cavity inverted cone	Class IIa
D002424001600	Steel burs cavity inverted cone	Class IIa
D002424001800	Steel burs cavity inverted cone	Class IIa
D002424002100	Steel burs cavity inverted cone	Class IIa
D002424002300	Steel burs cavity inverted cone	Class IIa
D002424002500	Steel burs cavity inverted cone	Class IIa
D002424002700	Steel burs cavity inverted cone	Class IIa
D002424002900	Steel burs cavity inverted cone	Class IIa
D002425000600	Steel burs cavity inverted cone	Class IIa
D002425000700	Steel burs cavity inverted cone	Class IIa
D002425000800	Steel burs cavity inverted cone	Class IIa
D002425000900	Steel burs cavity inverted cone	Class IIa
D002425001000	Steel burs cavity inverted cone	Class IIa
D002425001200	Steel burs cavity inverted cone	Class IIa
D002425001400	Steel burs cavity inverted cone	Class IIa
D002425001600	Steel burs cavity inverted cone	Class IIa
D002425001800	Steel burs cavity inverted cone	Class IIa
D002425002100	Steel burs cavity inverted cone	Class IIa

LNE(G-MED (0459) reconnaît que son certificat CE est  
 valide pour les dispositifs médicaux décrits  
 ..... 2015  
 LNE(G-MED (0459) recognizes that its EC certificate  
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Item number	Product designation	Product class
D002814000700	Steel burs cavity cylindric round	Class IIa
D002814000900	Steel burs cavity cylindric round	Class IIa
D002814001000	Steel burs cavity cylindric round	Class IIa
D002814001200	Steel burs cavity cylindric round	Class IIa
D002814001400	Steel burs cavity cylindric round	Class IIa
D002814001600	Steel burs cavity cylindric round	Class IIa
D002814001800	Steel burs cavity cylindric round	Class IIa
D002814002100	Steel burs cavity cylindric round	Class IIa
D002814002300	Steel burs cavity cylindric round	Class IIa
D002824000800	Steel burs cavity cylindric round	Class IIa
D002824000900	Steel burs cavity cylindric round	Class IIa
D002824001000	Steel burs cavity cylindric round	Class IIa
D002824001200	Steel burs cavity cylindric round	Class IIa
D002824001400	Steel burs cavity cylindric round	Class IIa
D002824001600	Steel burs cavity cylindric round	Class IIa
D002824001800	Steel burs cavity cylindric round	Class IIa
D002824002100	Steel burs cavity cylindric round	Class IIa
D002824002300	Steel burs cavity cylindric round	Class IIa
D003414000700	Steel burs cav. cyl. square	Class IIa
D003414000800	Steel burs cav. cyl. square	Class IIa
D003414000900	Steel burs cav. cyl. square	Class IIa
D003414001000	Steel burs cav. cyl. square	Class IIa
D003414001200	Steel burs cav. cyl. square	Class IIa
D003414001400	Steel burs cav. cyl. square	Class IIa
D003414001600	Steel burs cav. cyl. square	Class IIa
D003414001800	Steel burs cav. cyl. square	Class IIa
D003414002100	Steel burs cav. cyl. square	Class IIa
D003414002300	Steel burs cav. cyl. square	Class IIa
D003414002500	Steel burs cav. cyl. square	Class IIa
D003414002700	Steel burs cav. cyl. square	Class IIa
D003424000700	Steel burs cav. cyl. square	Class IIa
D003424000800	Steel burs cav. cyl. square	Class IIa
D003424000900	Steel burs cav. cyl. square	Class IIa
D003424001000	Steel burs cav. cyl. square	Class IIa
D003424001200	Steel burs cav. cyl. square	Class IIa
D003424001400	Steel burs cav. cyl. square	Class IIa
D003424001600	Steel burs cav. cyl. square	Class IIa
D003424001800	Steel burs cav. cyl. square	Class IIa
D003424002100	Steel burs cav. cyl. square	Class IIa
D003424002300	Steel burs cav. cyl. square	Class IIa
D003424002500	Steel burs cav. cyl. square	Class IIa
D003424002700	Steel burs cav. cyl. square	Class IIa
D003425000700	Steel burs cav. cyl. square	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est  
valide pour les dispositifs médicaux décrits  
...20-AOÛT-2015...  
LNE/G-MED (0459) recognizes that its EC certificate  
is valid for the medical devices listed

Item number	Product designation	Product class
D00342500800	Steel burs cav. cyl. square	Class IIa
D00342500900	Steel burs cav. cyl. square	Class IIa
D003425001000	Steel burs cav. cyl. square	Class IIa
D003425001200	Steel burs cav. cyl. square	Class IIa
D003425001400	Steel burs cav. cyl. square	Class IIa
D003425001600	Steel burs cav. cyl. square	Class IIa
D003425001800	Steel burs cav. cyl. square	Class IIa
D003425002100	Steel burs cav. cyl. square	Class IIa
D003425002300	Steel burs cav. cyl. square	Class IIa
D003814000700	Steel burs cav cone square	Class IIa
D003814000800	Steel burs cav cone square	Class IIa
D003814000900	Steel burs cav cone square	Class IIa
D003814001000	Steel burs cav cone square	Class IIa
D003814001200	Steel burs cav cone square	Class IIa
D003814001400	Steel burs cav cone square	Class IIa
D003814001600	Steel burs cav cone square	Class IIa
D003814001800	Steel burs cav cone square	Class IIa
D003814002100	Steel burs cav cone square	Class IIa
D003814002300	Steel burs cav cone square	Class IIa
D003814002500	Steel burs cav cone square	Class IIa
D003824000800	Steel burs cav cone square	Class IIa
D003824000900	Steel burs cav cone square	Class IIa
D003824001000	Steel burs cav cone square	Class IIa
D003824001200	Steel burs cav cone square	Class IIa
D003824001400	Steel burs cav cone square	Class IIa
D003824001600	Steel burs cav cone square	Class IIa
D003824001800	Steel burs cav cone square	Class IIa
D003824002100	Steel burs cav cone square	Class IIa
D003824002300	Steel burs cav cone square	Class IIa
D003824002500	Steel burs cav cone square	Class IIa
D003825000900	Steel burs cav cone square	Class IIa
D003825001000	Steel burs cav cone square	Class IIa
D003825001200	Steel burs cav cone square	Class IIa
D003825001400	Steel burs cav cone square	Class IIa
D003825001600	Steel burs cav cone square	Class IIa
D003825001800	Steel burs cav cone square	Class IIa
D003825002100	Steel burs cav cone square	Class IIa
D003825002300	Steel burs cav cone square	Class IIa
D004124001200	Batt	Class IIa
D004124001400	Batt	Class IIa
D004124001600	Batt	Class IIa
D004124001800	Batt	Class IIa
D004124002100	Batt	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits  
 20 AOÛT 2015  
 LNE/G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed

Item number	Product designation	Product class
D004124002300	Batt	Class IIa
D004125001200	Batt	Class IIa
D004125001400	Batt	Class IIa
D004125001600	Batt	Class IIa
D004125001800	Batt	Class IIa
D004125002100	Batt	Class IIa
D004125002300	Batt	Class IIa
D006114001000	Steel burs finishing round	Class IIa
D006114001200	Steel burs finishing round	Class IIa
D006114001400	Steel burs finishing round	Class IIa
D006114001600	Steel burs finishing round	Class IIa
D006114001800	Steel burs finishing round	Class IIa
D006114002100	Steel burs finishing round	Class IIa
D006114002300	Steel burs finishing round	Class IIa
D006114002500	Steel burs finishing round	Class IIa
D006114002700	Steel burs finishing round	Class IIa
D006114002900	Steel burs finishing round	Class IIa
D006114003500	Steel burs finishing round	Class IIa
D006114003700	Steel burs finishing round	Class IIa
D006114004000	Steel burs finishing round	Class IIa
D006124001000	Steel burs finishing round	Class IIa
D006124001200	Steel burs finishing round	Class IIa
D006124001400	Steel burs finishing round	Class IIa
D006124001600	Steel burs finishing round	Class IIa
D006124001800	Steel burs finishing round	Class IIa
D006124002100	Steel burs finishing round	Class IIa
D006124002300	Steel burs finishing round	Class IIa
D006124002500	Steel burs finishing round	Class IIa
D006124002700	Steel burs finishing round	Class IIa
D006124002900	Steel burs finishing round	Class IIa
D006124003100	Steel burs finishing round	Class IIa
D006124003300	Steel burs finishing round	Class IIa
D006124003500	Steel burs finishing round	Class IIa
D006124003700	Steel burs finishing round	Class IIa
D006124004000	Steel burs finishing round	Class IIa
D006214001000	Steel burs finishing oval	Class IIa
D006214001200	Steel burs finishing oval	Class IIa
D006214001400	Steel burs finishing oval	Class IIa
D006214001600	Steel burs finishing oval	Class IIa
D006214001800	Steel burs finishing oval	Class IIa
D006214002100	Steel burs finishing oval	Class IIa
D006214002300	Steel burs finishing oval	Class IIa
D006214002500	Steel burs finishing oval	Class IIa

LNE(G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits

LNE(G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed

2015

Item number	Product designation	Product class
D006214002700	Steel burs finishing oval	Class IIa
D006214002900	Steel burs finishing oval	Class IIa
D006224001000	Steel burs finishing oval	Class IIa
D006224001200	Steel burs finishing oval	Class IIa
D006224001400	Steel burs finishing oval	Class IIa
D006224001600	Steel burs finishing oval	Class IIa
D006224001800	Steel burs finishing oval	Class IIa
D006224002100	Steel burs finishing oval	Class IIa
D006224002300	Steel burs finishing oval	Class IIa
D006224002500	Steel burs finishing oval	Class IIa
D006224002900	Steel burs finishing oval	Class IIa
D006224003300	Steel burs finishing oval	Class IIa
D006314001000	Steel burs finishing pear	Class IIa
D006314001200	Steel burs finishing pear	Class IIa
D006314001400	Steel burs finishing pear	Class IIa
D006314001600	Steel burs finishing pear	Class IIa
D006314001800	Steel burs finishing pear	Class IIa
D006314002100	Steel burs finishing pear	Class IIa
D006314002300	Steel burs finishing pear	Class IIa
D006324001000	Steel burs finishing pear	Class IIa
D006324001200	Steel burs finishing pear	Class IIa
D006324001400	Steel burs finishing pear	Class IIa
D006324001600	Steel burs finishing pear	Class IIa
D006324001800	Steel burs finishing pear	Class IIa
D006324002100	Steel burs finishing pear	Class IIa
D006324002300	Steel burs finishing pear	Class IIa
D006324002500	Steel burs finishing pear	Class IIa
D006324002700	Steel burs finishing pear	Class IIa
D006324003100	Steel burs finishing pear	Class IIa
D006324003300	Steel burs finishing pear	Class IIa
D006324004000	Steel burs finishing pear	Class IIa
D006414001000	Steel burs finishing flame	Class IIa
D006414001200	Steel burs finishing flame	Class IIa
D006414001400	Steel burs finishing flame	Class IIa
D006414001600	Steel burs finishing flame	Class IIa
D006414001800	Steel burs finishing flame	Class IIa
D006414002100	Steel burs finishing flame	Class IIa
D006414002300	Steel burs finishing flame	Class IIa
D006414002500	Steel burs finishing flame	Class IIa
D006414002900	Steel burs finishing flame	Class IIa
D006414004000	Steel burs finishing flame	Class IIa
D006424001000	Steel burs finishing flame	Class IIa
D006424001200	Steel burs finishing flame	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est  
 valide pour les dispositifs médicaux décrits  
**20 AOÛT 2019**  
 LNE/G-MED (0459) recognizes that its EC certificate  
 is valid for the medical devices listed

Item number	Product designation	Product class
D006424001400	Steel burs finishing flame	Class IIa
D006424001600	Steel burs finishing flame	Class IIa
D006424001800	Steel burs finishing flame	Class IIa
D006424002100	Steel burs finishing flame	Class IIa
D006424002300	Steel burs finishing flame	Class IIa
D006424002500	Steel burs finishing flame	Class IIa
D006424002700	Steel burs finishing flame	Class IIa
D006424002900	Steel burs finishing flame	Class IIa
D006424003300	Steel burs finishing flame	Class IIa
D006424003500	Steel burs finishing flame	Class IIa
D006424003700	Steel burs finishing flame	Class IIa
D006424004000	Steel burs finishing flame	Class IIa
D020524000600	LN	Class IIa
D065B14001000	Steel burs finishing bud	Class IIa
D065B14001800	Steel burs finishing bud	Class IIa
D065B14002100	Steel burs finishing bud	Class IIa
D065B14002700	Steel burs finishing bud	Class IIa
D065B24001000	Steel burs finishing bud	Class IIa
D065B24001200	Steel burs finishing bud	Class IIa
D065B24001400	Steel burs finishing bud	Class IIa
D065B24001600	Steel burs finishing bud	Class IIa
D065B24001800	Steel burs finishing bud	Class IIa
D065B24002100	Steel burs finishing bud	Class IIa
D065B24002300	Steel burs finishing bud	Class IIa
D065B24002900	Steel burs finishing bud	Class IIa
D065B24004000	Steel burs finishing bud	Class IIa
D171I14001200	Surgical burs round, coarse	Class IIa
D171I14001400	Surgical burs round, coarse	Class IIa
D171I14001600	Surgical burs round, coarse	Class IIa
D171I14001800	Surgical burs round, coarse	Class IIa
D171I14002100	Surgical burs round, coarse	Class IIa
D171I14002300	Surgical burs round, coarse	Class IIa
D171I14002500	Surgical burs round, coarse	Class IIa
D171I14002700	Surgical burs round, coarse	Class IIa
D171I14003100	Surgical burs round, coarse	Class IIa
D171I14003300	Surgical burs round, coarse	Class IIa
D171I14003500	Surgical burs round, coarse	Class IIa
D171I14003700	Surgical burs round, coarse	Class IIa
D171I14004000	Surgical burs round, coarse	Class IIa
D171I14004500	Surgical burs round, coarse	Class IIa
D171I14005000	Surgical burs round, coarse	Class IIa
D171I14005500	Surgical burs round, coarse	Class IIa
D171I14006000	Surgical burs round, coarse	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est  
 valide pour les dispositifs médicaux décrits  
 ..... 23 AOÛT 2015 .....  
 LNE/G-MED (0459) recognizes that its EC certificate  
 is valid for the medical devices listed

Item number	Product designation	Product class
D171114006500	Surgical burs round, coarse	Class IIa
D172114002100	Surgical burs cone, coarse	Class IIa
D172114003300	Surgical burs cone, coarse	Class IIa
E012314000400	TC cavity burs round	Class IIa
E012314000500	TC cavity burs round	Class IIa
E012314000600	TC cavity burs round	Class IIa
E012314000700	TC cavity burs round	Class IIa
E012314000800	TC cavity burs round	Class IIa
E012314000900	TC cavity burs round	Class IIa
E012314001000	TC cavity burs round	Class IIa
E012314001200	TC cavity burs round	Class IIa
E012314001400	TC cavity burs round	Class IIa
E012314001600	TC cavity burs round	Class IIa
E012314001800	TC cavity burs round	Class IIa
E012314002100	TC cavity burs round	Class IIa
E012314002300	TC cavity burs round	Class IIa
E012314002500	TC cavity burs round	Class IIa
E012314002700	TC cavity burs round	Class IIa
E012324100500	TC cavity burs round	Class IIa
E012324100600	TC cavity burs round	Class IIa
E012324100700	TC cavity burs round	Class IIa
E012324100800	TC cavity burs round	Class IIa
E012324100900	TC cavity burs round	Class IIa
E012324101000	TC cavity burs round	Class IIa
E012324101200	TC cavity burs round	Class IIa
E012324101400	TC cavity burs round	Class IIa
E012324101600	TC cavity burs round	Class IIa
E012324101800	TC cavity burs round	Class IIa
E012324102100	TC cavity burs round	Class IIa
E012324102300	TC cavity burs round	Class IIa
E012324102700	TC cavity burs round	Class IIa
E012332100900	TC cavity burs round	Class IIa
E012332101400	TC cavity burs round	Class IIa
E012334100500	TC cavity burs round	Class IIa
E012334100600	TC cavity burs round	Class IIa
E012334100700	TC cavity burs round	Class IIa
E012334100800	TC cavity burs round	Class IIa
E012334100900	TC cavity burs round	Class IIa
E012334101000	TC cavity burs round	Class IIa
E012334101200	TC cavity burs round	Class IIa
E012334101400	TC cavity burs round	Class IIa
E012334101600	TC cavity burs round	Class IIa
E012334101800	TC cavity burs round	Class IIa

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 valide pour les dispositifs médicaux décrits  
 2015  
 LNE/G-MED (0459) recognizes that its EC certificate  
 is valid for the medical devices listed

Item number	Product designation	Product class
E012334102100	TC cavity burs round	Class IIa
E012334102300	TC cavity burs round	Class IIa
E012334102700	TC cavity burs round	Class IIa
E012337101000	TC cavity burs round	Class IIa
E012337101200	TC cavity burs round	Class IIa
E012337101400	TC cavity burs round	Class IIa
E012337101600	TC cavity burs round	Class IIa
E012337101800	TC cavity burs round	Class IIa
E012337102100	TC cavity burs round	Class IIa
E012337102300	TC cavity burs round	Class IIa
E012337102700	TC cavity burs round	Class IIa
E012414000800	TC cavity burs inverted cone	Class IIa
E012414000900	TC cavity burs inverted cone	Class IIa
E012414001000	TC cavity burs inverted cone	Class IIa
E012414001200	TC cavity burs inverted cone	Class IIa
E012414001400	TC cavity burs inverted cone	Class IIa
E012414001600	TC cavity burs inverted cone	Class IIa
E012414001800	TC cavity burs inverted cone	Class IIa
E012424100600	TC cavity burs inverted cone	Class IIa
E012424100700	TC cavity burs inverted cone	Class IIa
E012424100800	TC cavity burs inverted cone	Class IIa
E012424100900	TC cavity burs inverted cone	Class IIa
E012424101000	TC cavity burs inverted cone	Class IIa
E012424101200	TC cavity burs inverted cone	Class IIa
E012424101400	TC cavity burs inverted cone	Class IIa
E012424101600	TC cavity burs inverted cone	Class IIa
E012424101800	TC cavity burs inverted cone	Class IIa
E012432101000	TC cavity burs inverted cone	Class IIa
E012434100600	TC cavity burs inverted cone	Class IIa
E012434100700	TC cavity burs inverted cone	Class IIa
E012434100800	TC cavity burs inverted cone	Class IIa
E012434100900	TC cavity burs inverted cone	Class IIa
E012434101000	TC cavity burs inverted cone	Class IIa
E012434101200	TC cavity burs inverted cone	Class IIa
E012434101400	TC cavity burs inverted cone	Class IIa
E012434101600	TC cavity burs inverted cone	Class IIa
E012434101800	TC cavity burs inverted cone	Class IIa
E012814001000	TC cavity burs cylinder round	Class IIa
E012814001200	TC cavity burs cylinder round	Class IIa
E012814001400	TC cavity burs cylinder round	Class IIa
E012824100900	TC cavity burs cylinder round	Class IIa
E012824101000	TC cavity burs cylinder round	Class IIa
E012824101200	TC cavity burs cylinder round	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits ..... 20.001.2015 ..... LNE/G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed

Item number	Product designation	Product class
E012824101400	TC cavity burs cylinder round	Class IIa
E012832100800	TC cavity burs cylinder round	Class IIa
E012834100800	TC cavity burs cylinder round	Class IIa
E012834101000	TC cavity burs cylinder round	Class IIa
E012834101200	TC cavity burs cylinder round	Class IIa
E012834101400	TC cavity burs cylinder round	Class IIa
E012914000800	TC cavity burs cylinder square	Class IIa
E012914000900	TC cavity burs cylinder square	Class IIa
E012914001000	TC cavity burs cylinder square	Class IIa
E012914001200	TC cavity burs cylinder square	Class IIa
E012914001400	TC cavity burs cylinder square	Class IIa
E012914001600	TC cavity burs cylinder square	Class IIa
E012924100800	TC cavity burs cylinder square	Class IIa
E012924101000	TC cavity burs cylinder square	Class IIa
E012924101200	TC cavity burs cylinder square	Class IIa
E012924101400	TC cavity burs cylinder square	Class IIa
E012924101600	TC cavity burs cylinder square	Class IIa
E012934100800	TC cavity burs cylinder square	Class IIa
E012934100900	TC cavity burs cylinder square	Class IIa
E012934101000	TC cavity burs cylinder square	Class IIa
E012934101200	TC cavity burs cylinder square	Class IIa
E012934101400	TC cavity burs cylinder square	Class IIa
E012934101600	TC cavity burs cylinder square	Class IIa
E012937101000	TC cavity burs cylinder square	Class IIa
E012937101200	TC cavity burs cylinder square	Class IIa
E013014000900	TC cavity burs cone square	Class IIa
E013014001000	TC cavity burs cone square	Class IIa
E013014001200	TC cavity burs cone square	Class IIa
E013014001400	TC cavity burs cone square	Class IIa
E013014001600	TC cavity burs cone square	Class IIa
E013014001800	TC cavity burs cone square	Class IIa
E013024100800	TC cavity burs cone square	Class IIa
E013024100900	TC cavity burs cone square	Class IIa
E013024101000	TC cavity burs cone square	Class IIa
E013024101200	TC cavity burs cone square	Class IIa
E013024101400	TC cavity burs cone square	Class IIa
E013032101000	TC cavity burs cone square	Class IIa
E013034100800	TC cavity burs cone square	Class IIa
E013034100900	TC cavity burs cone square	Class IIa
E013034101000	TC cavity burs cone square	Class IIa
E013034101200	TC cavity burs cone square	Class IIa
E013034101400	TC cavity burs cone square	Class IIa
E013034101600	TC cavity burs cone square	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits

LNE/G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed

2015

Item number	Product designation	Product class
E013034101800	TC cavity burs cone square	Class IIa
E013037101000	TC cavity burs cone square	Class IIa
E013037101200	TC cavity burs cone square	Class IIa
E013614001000	TC cavity burs cylinder round	Class IIa
E013614001200	TC cavity burs cylinder round	Class IIa
E013614001400	TC cavity burs cylinder round	Class IIa
E013624100900	TC cavity burs cylinder round	Class IIa
E013624101000	TC cavity burs cylinder round	Class IIa
E013632100800	TC cavity burs cylinder round	Class IIa
E013634100800	TC cavity burs cylinder round	Class IIa
E013634101000	TC cavity burs cylinder round	Class IIa
E013634101200	TC cavity burs cylinder round	Class IIa
E013634101400	TC cavity burs cylinder round	Class IIa
E013714000800	TC cavity burs cylinder square	Class IIa
E013714000900	TC cavity burs cylinder square	Class IIa
E013714001000	TC cavity burs cylinder square	Class IIa
E013714001200	TC cavity burs cylinder square	Class IIa
E013714001400	TC cavity burs cylinder square	Class IIa
E013714001600	TC cavity burs cylinder square	Class IIa
E013724100800	TC cavity burs cylinder square	Class IIa
E013724101000	TC cavity burs cylinder square	Class IIa
E013724101200	TC cavity burs cylinder square	Class IIa
E013724101400	TC cavity burs cylinder square	Class IIa
E013724101800	TC cavity burs cylinder square	Class IIa
E013732101000	TC cavity burs cylinder square	Class IIa
E013734100800	TC cavity burs cylinder square	Class IIa
E013734100900	TC cavity burs cylinder square	Class IIa
E013734101000	TC cavity burs cylinder square	Class IIa
E013734101200	TC cavity burs cylinder square	Class IIa
E013734101400	TC cavity burs cylinder square	Class IIa
E013734101600	TC cavity burs cylinder square	Class IIa
E013734101800	TC cavity burs cylinder square	Class IIa
E013734102100	TC cavity burs cylinder square	Class IIa
E013737100800	TC cavity burs cylinder square	Class IIa
E013737101000	TC cavity burs cylinder square	Class IIa
E013737101200	TC cavity burs cylinder square	Class IIa
E013737101400	TC cavity burs cylinder square	Class IIa
E013814000800	TC cavity burs cone square	Class IIa
E013814000900	TC cavity burs cone square	Class IIa
E013814001200	TC cavity burs cone square	Class IIa
E013814001400	TC cavity burs cone square	Class IIa
E013814001600	TC cavity burs cone square	Class IIa
E013814001800	TC cavity burs cone square	Class IIa

LNE(G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits  
 20-AOUT-2015  
 LNE(G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed

Item number	Product designation	Product class
E013814002100	TC cavity burs cone square	Class IIa
E013814002300	TC cavity burs cone square	Class IIa
E013824100900	TC cavity burs cone square	Class IIa
E013824101000	TC cavity burs cone square	Class IIa
E013824101200	TC cavity burs cone square	Class IIa
E013824101400	TC cavity burs cone square	Class IIa
E013824101600	TC cavity burs cone square	Class IIa
E013824102100	TC cavity burs cone square	Class IIa
E013832101000	TC cavity burs cone square	Class IIa
E013834100800	TC cavity burs cone square	Class IIa
E013834100900	TC cavity burs cone square	Class IIa
E013834101000	TC cavity burs cone square	Class IIa
E013834101200	TC cavity burs cone square	Class IIa
E013834101400	TC cavity burs cone square	Class IIa
E013834101600	TC cavity burs cone square	Class IIa
E013834102100	TC cavity burs cone square	Class IIa
E013837101000	TC cavity burs cone square	Class IIa
E013837101200	TC cavity burs cone square	Class IIa
E013837101600	TC cavity burs cone square	Class IIa
E013837102100	TC cavity burs cone square	Class IIa
E014714000900	TC cavity burs cone round	Class IIa
E014734100900	TC cavity burs cone round	Class IIa
E014834101000	TC cavity burs cone round	Class IIa
E014834101200	TC cavity burs cone round	Class IIa
E014914001000	TC cavity burs cone round	Class IIa
E014914001200	TC cavity burs cone round	Class IIa
E014924101000	TC cavity burs cone round	Class IIa
E014924101400	TC cavity burs cone round	Class IIa
E014924101600	TC cavity burs cone round	Class IIa
E014934101000	TC cavity burs cone round	Class IIa
E014934101200	TC cavity burs cone round	Class IIa
E014934101400	TC cavity burs cone round	Class IIa
E014934101600	TC cavity burs cone round	Class IIa
E015114100000	Zekrya	Class IIa
E015115100000	Zekrya	Class IIa
E015124100000	Zekrya LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits	Class IIa
E015134100000	Zekrya	Class IIa
E015135100000	Zekrya LNE/G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed	Class IIa
E015224100000	Endo-Z	Class IIa
E015234100000	Endo-Z	Class IIa
E015237100000	Endo-Z	Class IIa
E015334101200	Transmetal	Class IIa
E015337101200	Transmetal	Class IIa

Item number	Product designation	Product class
E015434100900	Transmetal	Class IIa
E015434101000	Transmetal	Class IIa
E015434101200	Transmetal	Class IIa
E015437101000	Transmetal	Class IIa
E015534101000	DC	Class IIa
E015534101200	DC	Class IIa
E015634101000	DC	Class IIa
E015634101200	DC	Class IIa
E015734100000	Day-Z	Class IIa
E015834101000	DC	Class IIa
E015834101200	DC	Class IIa
E020524100610	LN	Class IIa
E020524100800	LN	Class IIa
E020524100810	LN	Class IIa
E020524101010	LN	Class IIa
E020524101210	LN	Class IIa
E020524101410	LN	Class IIa
E020524101610	LN	Class IIa
E020524190000	LN	Class IIa
E051034100800	CF	Class IIa
E051034101000	CF	Class IIa
E051034101400	CF	Class IIa
E051134101400	CF	Class IIa
E051134101600	CF	Class IIa
E051234101800	CF	Class IIa
E051234102300	CF	Class IIa
E051534101200	CF	Class IIa
E051634101400	CF	Class IIa
E051734100900	CF	Class IIa
E051734101000	CF	Class IIa
E051734101200	CF	Class IIa
E051834100700	CF	Class IIa
E051834100900	CF	Class IIa
E051834101000	CF	Class IIa
E051934101200	CF	Class IIa
E051934101400	CF	Class IIa
E051934101800	CF LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits	Class IIa
E052034101000	CF 2.0 AOÜT 2013	Class IIa
E052034101200	CF LNE/G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed	Class IIa
E052134101000	CF	Class IIa
E052134101200	CF	Class IIa
E052234100900	CF	Class IIa
E052334101200	CF	Class IIa

Item number	Product designation	Product class
E052434100900	CF	Class IIa
E052534101200	CF	Class IIa
E052634101000	CF	Class IIa
E052634101200	CF	Class IIa
E052734101400	CF	Class IIa
E052734101600	CF	Class IIa
E052834101000	CF	Class IIa
E052834101200	CF	Class IIa
E052834101400	CF	Class IIa
E052934101600	CF	Class IIa
E053034101400	CF	Class IIa
E053134101800	CF	Class IIa
E053234101200	CF	Class IIa
E053334101200	CF	Class IIa
E053334101400	CF	Class IIa
E053434101200	CF	Class IIa
E053434101400	CF	Class IIa
E053434101600	CF	Class IIa
E053434101800	CF	Class IIa
E053534101200	CF	Class IIa
E053634101400	CF	Class IIa
E053734101400	CF	Class IIa
E054014101600	CF	Class IIa
E054024101600	CF	Class IIa
E054035101600	CF	Class IIa
E054114101600	CF	Class IIa
E054124101600	CF	Class IIa
E054135101600	CF	Class IIa
E055034100100	CF	Class IIa
E055034102A00	CF	Class IIa
E055034102B00	CF	Class IIa
E055234101600	CF	Class IIa
E055334101600	CF	Class IIa
E056035100000	Aryane-Z	Class IIa
E057034101200	DC	Class IIa
E057134101000	DC	Class IIa
E057134101200	DC	Class IIa
E058034101000	Transmetal	Class IIa
E059004100000	Metal-Master	Class IIa
E123A24101000	Excavabur	Class IIa
E123A24101200	Excavabur	Class IIa
E123A24101400	Excavabur	Class IIa
E123A24101600	Excavabur	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits  
 20.08.2019  
 LNE/G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed

Item number	Product designation	Product class
E123A24101800	Excavabur	Class IIa
E123A24102100	Excavabur	Class IIa
E123A24102300	Excavabur	Class IIa
E123A24102500	Excavabur	Class IIa
E123A24102700	Excavabur	Class IIa
E123A34101000	Excavabur	Class IIa
E123A34101200	Excavabur	Class IIa
E123A34101400	Excavabur	Class IIa
E123A34101600	Excavabur	Class IIa
E123A34101800	Excavabur	Class IIa
E123A34102100	Excavabur	Class IIa
E123A34102300	Excavabur	Class IIa
E123A34102500	Excavabur	Class IIa
E123A34102700	Excavabur	Class IIa
E129X14101000	TCcavity burs cylinder square	Class IIa
E129X14101200	TCcavity burs cylinder square	Class IIa
E129X34101000	TCcavity burs cylinder square	Class IIa
E129X34101200	TCcavity burs cylinder square	Class IIa
E129X34101400	TCcavity burs cylinder square	Class IIa
E130L34100900	TC cavity burs cone square	Class IIa
E130L34101000	TC cavity burs cone square	Class IIa
E130X14101200	TC cavity burs cone square	Class IIa
E130X34101000	TC cavity burs cone square	Class IIa
E130X34101200	TC cavity burs cone square	Class IIa
E137X14101000	TC cavity burs cylinder square	Class IIa
E137X34101000	TC cavity burs cylinder square	Class IIa
E137X34101200	TC cavity burs cylinder square	Class IIa
E137X34101400	TC cavity burs cylinder square	Class IIa
E138L34100900	TC cavity burs cone square	Class IIa
E138L34101000	TC cavity burs cone square	Class IIa
E138L34101600	TC cavity burs cone square	Class IIa
E138X34101000	TC cavity burs cone square	Class IIa
E138X34101200	TC cavity burs cone square	Class IIa
E150A14000600	TC amalgam burs	Class IIa
E150A14000800	TC amalgam burs	Class IIa
E150A24100800	TC amalgam burs	Class IIa
E150A24101000	TC amalgam burs	Class IIa
E150A24101200	TC amalgam burs	Class IIa
E150A32100800	TC amalgam burs	Class IIa
E150A32101000	TC amalgam burs	Class IIa
E150A34100600	TC amalgam burs	Class IIa
E150A34100800	TC amalgam burs	Class IIa
E150A34100900	TC amalgam burs	Class IIa

Item number	Product designation	Product class
E150A34101000	TC amalgam burs	Class IIa
E150A34101200	TC amalgam burs	Class IIa
E150B24101000	TC amalgam burs	Class IIa
E150B24101200	TC amalgam burs	Class IIa
E150B24101400	TC amalgam burs	Class IIa
E150B34100800	TC amalgam burs	Class IIa
E150B34101000	TC amalgam burs	Class IIa
E150B34101200	TC amalgam burs	Class IIa
E150B34101400	TC amalgam burs	Class IIa
E150C34100800	TC amalgam burs	Class IIa
E150C34101400	TC amalgam burs	Class IIa
E510L34101400	CF	Class IIa
E536L34101400	CF LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits	Class IIa
E537L34101400	CF	Class IIa
E541A14101600	CF LNE/G-MED (0459) recognizes that its EC certificate	Class IIa
E541A24101600	CF is valid for the medical devices listed	Class IIa
E541A35101600	CF	Class IIa
F000133201000	Diamond burs	Class IIa
F000133201400	Diamond burs	Class IIa
F000134102300	Diamond burs	Class IIa
F000134200900	Diamond burs	Class IIa
F000134201000	Diamond burs	Class IIa
F000134201200	Diamond burs	Class IIa
F000134201400	Diamond burs	Class IIa
F000134201600	Diamond burs	Class IIa
F000134201800	Diamond burs	Class IIa
F000134202300	Diamond burs	Class IIa
F000134202500	Diamond burs	Class IIa
F000134203300	Diamond burs	Class IIa
F000134301000	Diamond burs	Class IIa
F000134301400	Diamond burs	Class IIa
F000134301600	Diamond burs	Class IIa
F000134301800	Diamond burs	Class IIa
F000134302300	Diamond burs	Class IIa
F000134401200	Diamond burs	Class IIa
F000134401400	Diamond burs	Class IIa
F000134401600	Diamond burs	Class IIa
F000134401800	Diamond burs	Class IIa
F000134402300	Diamond burs	Class IIa
F000234200900	Diamond burs	Class IIa
F000234201000	Diamond burs	Class IIa
F000234201200	Diamond burs	Class IIa
F000234301200	Diamond burs	Class IIa

Item number	Product designation	Product class
F000234301400	Diamond burs	Class IIa
F001034200900	Diamond burs	Class IIa
F001034201000	Diamond burs	Class IIa
F001034201200	Diamond burs	Class IIa
F001034201400	Diamond burs	Class IIa
F001034201600	Diamond burs	Class IIa
F001034201800	Diamond burs	Class IIa
F001034301400	Diamond burs	Class IIa
F001934201200	Diamond burs	Class IIa
F001934301000	Diamond burs	Class IIa
F001934301200	Diamond burs	Class IIa
F001934301400	Diamond burs	Class IIa
F003234201400	Diamond burs	Class IIa
F003234201600	Diamond burs	Class IIa
F003334202700	Diamond burs	Class IIa
F003834203300	Diamond burs	Class IIa
F003934203700	Diamond burs	Class IIa
F004134204500	Diamond burs	Class IIa
F004234204500	Diamond burs	Class IIa
F006834204000	Diamond burs	Class IIa
F006834304000	Diamond burs	Class IIa
F010733201000	Diamond burs	Class IIa
F010733201200	Diamond burs	Class IIa
F010734200800	Diamond burs	Class IIa
F010734200900	Diamond burs	Class IIa
F010734201000	Diamond burs	Class IIa
F010734201200	Diamond burs	Class IIa
F010734201400	Diamond burs	Class IIa
F010734201600	Diamond burs	Class IIa
F010734300900	Diamond burs	Class IIa
F010734301000	Diamond burs	Class IIa
F010734301200	Diamond burs	Class IIa
F010734301400	Diamond burs	Class IIa
F010734401000	Diamond burs	Class IIa
F010734401400	Diamond burs	Class IIa
F011034001000	Diamond burs	Class IIa
F011034201000	Diamond burs	Class IIa
F011034201200	Diamond burs	Class IIa
F011034201400	Diamond burs	Class IIa
F011034301200	Diamond burs	Class IIa
F011034301400	Diamond burs	Class IIa
F011034301600	Diamond burs	Class IIa
F011034301800	Diamond burs	Class IIa

LNE(G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits  
LNE(G-MED(0459) recognizes that its EC certificate is valid for the medical devices listed

Item number	Product designation	Product class
F011034401400	Diamond burs	Class IIa
F011134001000	Diamond burs	Class IIa
F011134201200	Diamond burs	Class IIa
F011134201400	Diamond burs	Class IIa
F011134301200	Diamond burs	Class IIa
F011134301400	Diamond burs	Class IIa
F011134301600	Diamond burs	Class IIa
F011134401400	Diamond burs	Class IIa
F011234001000	Diamond burs	Class IIa
F011234201400	Diamond burs	Class IIa
F012934201200	Diamond burs	Class IIa
F013034301200	Diamond burs	Class IIa
F013034301400	Diamond burs	Class IIa
F013134301400	Diamond burs	Class IIa
F013134301600	Diamond burs	Class IIa
F013734301200	Diamond burs	Class IIa
F014034001000	Diamond burs	Class IIa
F014034201400	Diamond burs	Class IIa
F014134001000	Diamond burs	Class IIa
F014134201200	Diamond burs	Class IIa
F014134201400	Diamond burs	Class IIa
F014134301200	Diamond burs	Class IIa
F014134301400	Diamond burs	Class IIa
F014234001000	Diamond burs	Class IIa
F014234201400	Diamond burs	Class IIa
F016534001400	Diamond burs	Class IIa
F016534101400	Diamond burs	Class IIa
F016534201200	Diamond burs	Class IIa
F016534201400	Diamond burs	Class IIa
F016534201600	Diamond burs	Class IIa
F016534301400	Diamond burs	Class IIa
F016534401400	Diamond burs	Class IIa
F016634001400	Diamond burs	Class IIa
F016634101000	Diamond burs	Class IIa
F016634201000	Diamond burs	Class IIa
F016634201400	Diamond burs	Class IIa
F016634201800	Diamond burs	Class IIa
F016634301400	Diamond burs	Class IIa
F016634301800	Diamond burs	Class IIa
F016734201000	Diamond burs	Class IIa
F016834201000	Diamond burs	Class IIa
F016834201200	Diamond burs	Class IIa
F016834201400	Diamond burs	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est valide pour les dispositifs médicaux décrits ..... 20-AQU-2015 ..... LNE/G-MED (0459) recognizes that its EC certificate is valid for the medical devices listed

Item number	Product designation	Product class
F016834201600	Diamond burs	Class IIa
F016834201800	Diamond burs	Class IIa
F016834202300	Diamond burs	Class IIa
F016834202500	Diamond burs	Class IIa
F017134101200	Diamond burs	Class IIa
F017134201200	Diamond burs	Class IIa
F017134201400	Diamond burs	Class IIa
F017134301200	Diamond burs	Class IIa
F017234201200	Diamond burs	Class IIa
F017234201400	Diamond burs	Class IIa
F017234201600	Diamond burs	Class IIa
F017234301600	Diamond burs	Class IIa
F017234301800	Diamond burs	Class IIa
F017234401400	Diamond burs	Class IIa
F017234401600	Diamond burs	Class IIa
F017334101800	Diamond burs	Class IIa
F017334201400	Diamond burs	Class IIa
F017334201600	Diamond burs	Class IIa
F017334201800	Diamond burs	Class IIa
F017334301800	Diamond burs	Class IIa
F017334401600	Diamond burs	Class IIa
F019734200900	Diamond burs	Class IIa
F019734202500	Diamond burs	Class IIa
F019734301200	Diamond burs	Class IIa
F019734301600	Diamond burs	Class IIa
F019734301800	Diamond burs	Class IIa
F019734402500	Diamond burs	Class IIa
F019834101600	Diamond burs	Class IIa
F019834101800	Diamond burs	Class IIa
F019834102300	Diamond burs	Class IIa
F019834201600	Diamond burs	Class IIa
F019834301200	Diamond burs	Class IIa
F019834301400	Diamond burs	Class IIa
F019834301600	Diamond burs	Class IIa
F019834301800	Diamond burs	Class IIa
F019834302300	Diamond burs	Class IIa
F019834401400	Diamond burs	Class IIa
F019834401600	Diamond burs	Class IIa
F019834401800	Diamond burs	Class IIa
F019934201200	Diamond burs	Class IIa
F019934201400	Diamond burs	Class IIa
F019934201600	Diamond burs	Class IIa
F019934201800	Diamond burs	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est  
 valide pour les dispositifs médicaux décrits  
 ..... 2 ..... 0407 2015  
 LNE/G-MED (0459) recognizes that its EC certificate  
 is valid for the medical devices listed

Item number	Product designation	Product class
F019934202300	Diamond burs	Class IIa
F019934301400	Diamond burs	Class IIa
F019934401600	Diamond burs	Class IIa
F019934401800	Diamond burs	Class IIa
F019937301600	Diamond burs	Class IIa
F021934101200	Diamond burs	Class IIa
F021934201200	Diamond burs	Class IIa
F021934201600	Diamond burs	Class IIa
F022034201400	Diamond burs	Class IIa
F022534201200	Diamond burs	Class IIa
F022534201600	Diamond burs	Class IIa
F023334200900	Diamond burs	Class IIa
F023334201000	Diamond burs	Class IIa
F023334201200	Diamond burs	Class IIa
F023334201400	Diamond burs	Class IIa
F023334401200	Diamond burs	Class IIa
F023434201200	Diamond burs	Class IIa
F023434301200	Diamond burs	Class IIa
F023434301400	Diamond burs	Class IIa
F023434301600	Diamond burs	Class IIa
F023434301800	Diamond burs	Class IIa
F024534301000	Diamond burs	Class IIa
F024534301400	Diamond burs	Class IIa
F024933201200	Diamond burs	Class IIa
F024934001200	Diamond burs	Class IIa
F024934001400	Diamond burs	Class IIa
F024934001600	Diamond burs	Class IIa
F024934101200	Diamond burs	Class IIa
F024934101400	Diamond burs	Class IIa
F024934201000	Diamond burs	Class IIa
F024934301200	Diamond burs	Class IIa
F024934301400	Diamond burs	Class IIa
F024934301600	Diamond burs	Class IIa
F024934401200	Diamond burs	Class IIa
F024934401400	Diamond burs	Class IIa
F025034001200	Diamond burs	Class IIa
F025034101200	Diamond burs	Class IIa
F025034101600	Diamond burs	Class IIa
F025034201200	Diamond burs	Class IIa
F025034301200	Diamond burs	Class IIa
F025034301600	Diamond burs	Class IIa
F025734001600	Diamond burs	Class IIa
F025734002300	Diamond burs	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est  
 valide pour les dispositifs médicaux décrits  
 LNE/G-MED (0459) recognizes that its EC certificate  
 is valid for the medical devices listed

Item number	Product designation	Product class
F025734101600	Diamond burs	Class IIa
F025734102300	Diamond burs	Class IIa
F025734301600	Diamond burs	Class IIa
F025734302300	Diamond burs	Class IIa
F027734002300	Diamond burs	Class IIa
F027734101800	Diamond burs	Class IIa
F027734201800	Diamond burs	Class IIa
F027734202300	Diamond burs	Class IIa
F027734302300	Diamond burs	Class IIa
F028834001000	Diamond burs	Class IIa
F028834200900	Diamond burs	Class IIa
F028834201000	Diamond burs	Class IIa
F028834201200	Diamond burs	Class IIa
F028834201400	Diamond burs	Class IIa
F028834301000	Diamond burs	Class IIa
F028834301200	Diamond burs	Class IIa
F028934001000	Diamond burs	Class IIa
F028934101400	Diamond burs	Class IIa
F028934201000	Diamond burs	Class IIa
F028934201200	Diamond burs	Class IIa
F028934201400	Diamond burs	Class IIa
F028934301200	Diamond burs	Class IIa
F028934301400	Diamond burs	Class IIa
F028934301600	Diamond burs	Class IIa
F028934401400	Diamond burs	Class IIa
F029034001000	Diamond burs	Class IIa
F029034101200	Diamond burs	Class IIa
F029034101400	Diamond burs	Class IIa
F029034201400	Diamond burs	Class IIa
F029034301200	Diamond burs	Class IIa
F029034301400	Diamond burs	Class IIa
F029034301600	Diamond burs	Class IIa
F029734301200	Diamond burs	Class IIa
F029734301400	Diamond burs	Class IIa
F029734301600	Diamond burs	Class IIa
F029834101600	Diamond burs	Class IIa
F029834201200	Diamond burs	Class IIa
F029834201600	Diamond burs	Class IIa
F029834301200	Diamond burs	Class IIa
F029834301400	Diamond burs	Class IIa
F029834301600	Diamond burs	Class IIa
F029834301800	Diamond burs	Class IIa
F029834302100	Diamond burs	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est  
 valide pour les dispositifs médicaux décrits  
 ..... 23-AOÛT 2015 .....  
 LNE/G-MED (0459) recognizes that its EC certificate  
 is valid for the medical devices listed

Item number	Product designation	Product class
F029834401600	Diamond burs	Class IIa
F029934101600	Diamond burs	Class IIa
F029934101800	Diamond burs	Class IIa
F029934201600	Diamond burs	Class IIa
F029934301200	Diamond burs	Class IIa
F029934301400	Diamond burs	Class IIa
F029934301600	Diamond burs	Class IIa
F029934301800	Diamond burs	Class IIa
F029934302100	Diamond burs	Class IIa
F029934401600	Diamond burs	Class IIa
F030434201600	Diamond burs	Class IIa
F030434202300	Diamond burs	Class IIa
F049634201200	Diamond burs	Class IIa
F054033200900	Diamond burs	Class IIa
F054034100900	Diamond burs	Class IIa
F054034101000	Diamond burs	Class IIa
F054034200900	Diamond burs	Class IIa
F054034300900	Diamond burs	Class IIa
F054034301000	Diamond burs	Class IIa
F054434201600	Diamond burs	Class IIa
F054534201600	Diamond burs	Class IIa
F054634301600	Diamond burs	Class IIa
F054634301800	Diamond burs	Class IIa
F069735301600	Diamond burs	Class IIa
F069735301800	Diamond burs	Class IIa
F095100000300	Isometrix burs	Class IIa
F095100000400	Isometrix burs	Class IIa
F095134000300	Isometrix burs	Class IIa
F095134000400	Isometrix burs	Class IIa
F095134090000	Isometrix burs	Class IIa
F095134100100	Isometrix burs	Class IIa
F095134100200	Isometrix burs	Class IIa
F095134100300	Isometrix burs	Class IIa
F095134100400	Isometrix burs	Class IIa
F095134190000	Isometrix burs	Class IIa
F095134200100	Isometrix burs	Class IIa
F095134200200	Isometrix burs	Class IIa
F095134200300	Isometrix burs	Class IIa
F095134200400	Isometrix burs	Class IIa
F095230090000	Isometrix burs	Class IIa
F095234000100	Isometrix burs	Class IIa
F095234000200	Isometrix burs	Class IIa
F095333001400	Diamond burs	Class IIa

LNE/G-MED (0459) reconnaît que son certificat CE est  
 valide pour les dispositifs médicaux décrits  
 20.01.2015  
 LNE/G-MED (0459) recognizes that its EC certificate  
 is valid for the medical devices listed

Item number	Product designation	Product class
F095334001400	Diamond burs	Class IIa
F095335001400	Diamond burs	Class IIa
F116100000000	Gingikit set	Class IIa
F116134210100	Gingikit set	Class IIa
F116134210200	Gingikit set	Class IIa
F116134210300	Gingikit set	Class IIa
F116135210400	Gingikit set	Class IIa
F116135210500	Gingikit set	Class IIa
F116135210600	Gingikit set	Class IIa
F116136210700	Gingikit set	Class IIa
F116136210800	Gingikit set	Class IIa
F116136210900	Gingikit set	Class IIa
F116200000000	Ceramprep set	Class IIa
F130000000000	Bogopolsky set	Class IIa
S209000310100	X-Post	Class IIa
S209000310200	X-Post	Class IIa
S209000310300	X-Post	Class IIa
S209000310400	X-Post	Class IIa

**Ballaigues, 5 August 2015**

**MAILLEFER INSTRUMENTS  
HOLDING Sàrl**



*Luc Etienne*  
**Luc Etienne**  
Management Representative

LNE/G-MED (0459) reconnaît que son certificat CE est  
valide pour les dispositifs médicaux décrits  
.....  
LNE/G-MED (0459) recognizes that its EC certificate  
is valid for the medical devices listed

**20 AOUT 2015**

# SERTIFIKATAS Nr. 10468<sub>4</sub> Peržiūrėjimas

Išleista 2015-08-20, Paryžius

Notifikavimo štaigos Nr.0459

## CE SERTIFIKATAS

Pilnas kokybės draudimo sistemos patvirtinimas  
93/42/EEB direktyva, II PRIEDO 4 dalis apimanti medicininės priemonės

Gamintojas

**MAILLEFER INSTRUMENTS HOLDING SARL**  
Chemin du Verger 3  
1338 BALLAIGUES SWITZERLAND  
(ŠVEICARIJA)

Priemonės(-ių) kategorija

**Sterilūs ir nesterilūs odontologiniai instrumentai, motorai ir kampiniai antgaliai, odontologinės rekonstrukcijos sistemos, odontologiniai prietaisai diagnozavimams**

LNE/G-MED patvirtina, kad pagal atliktus rezultatus, kurie nurodyti dokumente P129078, kokybės sistema taikoma dizainui, gamybai ir galutiniam patikrinimui medicinos priemonių, išvardintų sąraše viršuje, atitinka 93/42/EEB direktyvos, II priedo 4 dalies reikalavimus.

**Įsigaliojimo data: 2015-08-20**

**Galiojimo data: 2018-07-08**

LNE – 10468 4 perž.

**Cecile VAUGELADE**

**G-MED sertifikacijos skyriaus direktorius**



VERTIMAS TEISINGAS

Andrius Preišegalavičius

### *Prietaisų identifikacija*

Prietaisai, kuriems taikomas šis sertifikatas yra išvardinti gamintojo produktų sąrašė (55 puslapiai) autentifikuoto 2015 liepos 7 d. LNE/G-MED.

### *Identifikavimas vietos ir veiklos*

- Chemin du Verger, 3, 1338 BALLAIGUES, SUISSE:  
Kūrimas, gamyba, galutinis patikrinimas
- Chemin de Beau-Site, 5, 1338 BALLAIGUES, SUISSE:  
Gamyba
- Le creux, 1338 BALLAIGUES, SUISSE:  
Gamyba

VERTIMAS TEISINGAS

Direktorius

Andrius Preišegalavičius

LNE/G-MED	0459
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Cecile VAUGELADE  
G-MED sertifikacijos skyriaus  
direktorius



## EB Sertifikate Nr. 10468 4 perž. taikomų produktų sąrašas

Prod.nr.	Produkto žymėjimas	Produkto klasė
60667332	X-Post (kaištis)	II a klasė
60667333	X-Post (kaištis)	II a klasė
60667334	X-Post (kaištis)	II a klasė
60667335	X-Post (kaištis)	II a klasė
A000724000100	Šaknų kanalų grąžtas	II a klasė
A000724000200	Šaknų kanalų grąžtas	II a klasė
A000724000300	Šaknų kanalų grąžtas	II a klasė
A000724000400	Šaknų kanalų grąžtas	II a klasė
A000724090000	Šaknų kanalų grąžtas	II a klasė
A000823000100	Gates glidden grąžtas	II a klasė
A000823000103	Gates sterilus glidden grąžtas	II a klasė
A000823000200	Gates glidden grąžtas	II a klasė
A000823000203	Gates sterilus glidden grąžtas	II a klasė
A000823000300	Gates glidden grąžtas	II a klasė
A000823000303	Gates sterilus glidden grąžtas	II a klasė
A000823000400	Gates glidden grąžtas	II a klasė
A000823000403	Gates sterilus glidden grąžtas	II a klasė
A000823000500	Gates glidden grąžtas	II a klasė
A000823000503	Gates sterilus glidden grąžtas	II a klasė
A000823000600	Gates glidden grąžtas	II a klasė
A000823000603	Gates sterilus glidden grąžtas	II a klasė
A000823090100	Gates glidden grąžtas	II a klasė
A000823090103	Gates sterilus glidden grąžtas	II a klasė
A000824000100	Gates glidden grąžtas	II a klasė
A000824000103	Gates sterilus glidden grąžtas	II a klasė
A000824000200	Gates glidden grąžtas	II a klasė
A000824000203	Gates sterilus glidden grąžtas	II a klasė
A000824000300	Gates glidden grąžtas	II a klasė
A000824000303	Gates sterilus glidden grąžtas	II a klasė
A000824000400	Gates glidden grąžtas	II a klasė
A000824000403	Gates sterilus glidden grąžtas	II a klasė
A000824000500	Gates glidden grąžtas	II a klasė
A000824000503	Gates sterilus glidden grąžtas	II a klasė
A000824000600	Gates glidden grąžtas	II a klasė
A000824000603	Gates sterilus glidden grąžtas	II a klasė
A000824000700	Gates glidden grąžtas	II a klasė
A000824000703	Gates sterilus glidden grąžtas	II a klasė
A000824090100	Gates glidden grąžtas	II a klasė
A000824090103	Gates sterilus glidden grąžtas	II a klasė
A000825000100	Gates glidden grąžtas	II a klasė
A000825000103	Gates sterilus glidden grąžtas	II a klasė
A000825000200	Gates glidden grąžtas	II a klasė
A000825000203	Gates sterilus glidden grąžtas	II a klasė
A000825000300	Gates glidden grąžtas	II a klasė
A000825000303	Gates sterilus glidden grąžtas	II a klasė
A000825000400	Gates glidden grąžtas	II a klasė
A000825000403	Gates sterilus glidden grąžtas	II a klasė
A000825000500	Gates glidden grąžtas	II a klasė
A000825000503	Gates sterilus glidden grąžtas	II a klasė
A000825000600	Gates glidden grąžtas	II a klasė

<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
A000825000603	Gates sterilus glidden grąžtas	II a klasė
A000825090100	Gates glidden grąžtas	II a klasė
A000825090103	Gates sterilus glidden grąžtas	II a klasė
A000923000100	Largo Peeso Reamer	II a klasė
A000923000103	Largo sterilus	II a klasė
A000923000200	Largo Peeso Reamer	II a klasė
A000923000203	Largo sterilus	II a klasė
A000923000300	Largo Peeso Reamer	II a klasė
A000923000303	Largo sterilus	II a klasė
A000923000400	Largo Peeso Reamer	II a klasė
A000923000403	Largo sterilus	II a klasė
A000923000500	Largo Peeso Reamer	II a klasė
A000923000503	Largo sterilus	II a klasė
A000923000600	Largo Peeso Reamer	II a klasė
A000923000603	Largo sterilus	II a klasė
A000923090100	Largo Peeso Reamer	II a klasė
A000923090103	Largo sterilus	II a klasė
A000924000100	Largo Peeso Reamer	II a klasė
A000924000103	Largo sterilus	II a klasė
A000924000200	Largo Peeso Reamer	II a klasė
A000924000203	Largo sterilus	II a klasė
A000924000300	Largo Peeso Reamer	II a klasė
A000924000303	Largo sterilus	II a klasė
A000924000400	Largo Peeso Reamer	II a klasė
A000924000403	Largo sterilus	II a klasė
A000924000500	Largo Peeso Reamer	II a klasė
A000924000503	Largo sterilus	II a klasė
A000924000600	Largo Peeso Reamer	II a klasė
A000924000603	Largo sterilus	II a klasė
A000924090100	Largo Peeso Reamer	II a klasė
A000924090103	Largo sterilus	II a klasė
A001012501000	K-Reamer Torpan	II a klasė
A001012501500	K-Reamer Torpan	II a klasė
A001012502000	K-Reamer Torpan	II a klasė
A001012502500	K-Reamer Torpan	II a klasė
A001012503000	K-Reamer Torpan	II a klasė
A001012503500	K-Reamer Torpan	II a klasė
A001012504000	K-Reamer Torpan	II a klasė
A001012504500	K-Reamer Torpan	II a klasė
A001012505000	K-Reamer Torpan	II a klasė
A001012505500	K-Reamer Torpan	II a klasė
A001012506000	K-Reamer Torpan	II a klasė
A001012507000	K-Reamer Torpan	II a klasė
A001012508000	K-Reamer Torpan	II a klasė
A001012509000	K-Reamer Torpan	II a klasė
A001012510000	K-Reamer Torpan	II a klasė
A001012512000	K-Reamer Torpan	II a klasė
A001012590000	K-Reamer Torpan	II a klasė
A001021801000	K-Reamer Torpan	II a klasė
A001021801500	K-Reamer Torpan	II a klasė
A001021802000	K-Reamer Torpan	II a klasė
A001021802500	K-Reamer Torpan	II a klasė
A001021803000	K-Reamer Torpan	II a klasė
A001021803500	K-Reamer Torpan	II a klasė

<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
A001021804000	K-Reamer Torpan	II a klasė
A001021804500	K-Reamer Torpan	II a klasė
A001021805000	K-Reamer Torpan	II a klasė
A001021805500	K-Reamer Torpan	II a klasė
A001021806000	K-Reamer Torpan	II a klasė
A001021807000	K-Reamer Torpan	II a klasė
A001021808000	K-Reamer Torpan	II a klasė
A001021809000	K-Reamer Torpan	II a klasė
A001021810000	K-Reamer Torpan	II a klasė
A001021811000	K-Reamer Torpan	II a klasė
A001021812000	K-Reamer Torpan	II a klasė
A001021890000	K-Reamer Torpan	II a klasė
A001022101000	K-Reamer Torpan	II a klasė
A001022101500	K-Reamer Torpan	II a klasė
A001022102000	K-Reamer Torpan	II a klasė
A001022102500	K-Reamer Torpan	II a klasė
A001022103000	K-Reamer Torpan	II a klasė
A001022103500	K-Reamer Torpan	II a klasė
A001022104000	K-Reamer Torpan	II a klasė
A001022104500	K-Reamer Torpan	II a klasė
A001022105000	K-Reamer Torpan	II a klasė
A001022105500	K-Reamer Torpan	II a klasė
A001022106000	K-Reamer Torpan	II a klasė
A001022107000	K-Reamer Torpan	II a klasė
A001022108000	K-Reamer Torpan	II a klasė
A001022109000	K-Reamer Torpan	II a klasė
A001022110000	K-Reamer Torpan	II a klasė
A001022111000	K-Reamer Torpan	II a klasė
A001022112000	K-Reamer Torpan	II a klasė
A001022190000	K-Reamer Torpan	II a klasė
A001022190100	K-Reamer Torpan	II a klasė
A001022501000	K-Reamer Torpan	II a klasė
A001022501500	K-Reamer Torpan	II a klasė
A001022502000	K-Reamer Torpan	II a klasė
A001022502500	K-Reamer Torpan	II a klasė
A001022503000	K-Reamer Torpan	II a klasė
A001022503500	K-Reamer Torpan	II a klasė
A001022504000	K-Reamer Torpan	II a klasė
A001022504500	K-Reamer Torpan	II a klasė
A001022505000	K-Reamer Torpan	II a klasė
A001022505500	K-Reamer Torpan	II a klasė
A001022506000	K-Reamer Torpan	II a klasė
A001022507000	K-Reamer Torpan	II a klasė
A001022508000	K-Reamer Torpan	II a klasė
A001022509000	K-Reamer Torpan	II a klasė
A001022510000	K-Reamer Torpan	II a klasė
A001022511000	K-Reamer Torpan	II a klasė
A001022512000	K-Reamer Torpan	II a klasė
A001022590000	K-Reamer Torpan	II a klasė
A001022590100	K-Reamer Torpan	II a klasė
A001522101300	PATHFILE	II a klasė
A001522101303	PATHFILE STERILUS	II a klasė
A001522101600	PATHFILE	II a klasė
A001522101603	PATHFILE STERILUS	II a klasė
A001522101900	PATHFILE	II a klasė
A001522101903	PATHFILE STERILUS	II a klasė
A001522190000	PATHFILE	II a klasė

<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
A001522190003	PATHFILE STERILUS	II a klasė
A001522501300	PATHFILE	II a klasė
A001522501303	PATHFILE STERILUS	II a klasė
A001522501600	PATHFILE	II a klasė
A001522501603	PATHFILE STERILUS	II a klasė
A001522501900	PATHFILE	II a klasė
A001522501903	PATHFILE STERILUS	II a klasė
A001522590000	PATHFILE	II a klasė
A001522590003	PATHFILE STERILUS	II a klasė
A001522590200	PATHFILE	II a klasė
A001523101300	PATHFILE	II a klasė
A001523101303	PATHFILE STERILUS	II a klasė
A001523101600	PATHFILE	II a klasė
A001523101603	PATHFILE STERILUS	II a klasė
A001523101900	PATHFILE	II a klasė
A001523101903	PATHFILE STERILUS	II a klasė
A001523190000	PATHFILE	II a klasė
A001523190003	PATHFILE STERILUS	II a klasė
A002220090000	Lentulo pastos nešėjas (carrier)	II a klasė
A002221700100	Lentulo pastos nešėjas (carrier)	II a klasė
A002221700103	Lentulo sterilus pastos nešėjas (carrier)	II a klasė
A002221700200	Lentulo pastos nešėjas (carrier)	II a klasė
A002221700203	Lentulo sterilus pastos nešėjas (carrier)	II a klasė
A002221700300	Lentulo pastos nešėjas (carrier)	II a klasė
A002221700303	Lentulo sterilus pastos nešėjas (carrier)	II a klasė
A002221700400	Lentulo pastos nešėjas (carrier)	II a klasė
A002221700403	Lentulo sterilus pastos nešėjas (carrier)	II a klasė
A002221790000	Lentulo pastos nešėjas (carrier)	II a klasė
A002221790100	Lentulo pastos nešėjas (carrier)	II a klasė
A002222100100	Lentulo pastos nešėjas (carrier)	II a klasė
A002222100103	Lentulo sterilus pastos nešėjas (carrier)	II a klasė
A002222100200	Lentulo pastos nešėjas (carrier)	II a klasė
A002222100203	Lentulo sterilus pastos nešėjas (carrier)	II a klasė
A002222100300	Lentulo pastos nešėjas (carrier)	II a klasė
A002222100303	Lentulo sterilus pastos nešėjas (carrier)	II a klasė
A002222100400	Lentulo pastos nešėjas (carrier)	II a klasė
A002222100403	Lentulo sterilus pastos nešėjas (carrier)	II a klasė
A002222190000	Lentulo pastos nešėjas (carrier)	II a klasė
A002222190003	Lentulo sterilus pastos nešėjas (carrier)	II a klasė
A002222190100	Lentulo pastos nešėjas (carrier)	II a klasė
A002222500100	Lentulo pastos nešėjas (carrier)	II a klasė
A002222500103	Lentulo sterilus pastos nešėjas (carrier)	II a klasė
A002222500200	Lentulo pastos nešėjas (carrier)	II a klasė
A002222500203	Lentulo sterilus pastos nešėjas (carrier)	II a klasė
A002222500300	Lentulo pastos nešėjas (carrier)	II a klasė
A002222500303	Lentulo sterilus pastos nešėjas (carrier)	II a klasė
A002222500400	Lentulo pastos nešėjas (carrier)	II a klasė
A002222500403	Lentulo sterilus pastos nešėjas (carrier)	II a klasė
A002222590000	Lentulo pastos nešėjas (carrier)	II a klasė
A002222590003	Lentulo sterilus pastos nešėjas (carrier)	II a klasė
A0022225190100	Lentulo pastos nešėjas (carrier)	II a klasė
A004B02102001	Spiro COLORINOX	II a klasė
A004B02102501	Spiro COLORINOX	II a klasė
A004B02103001	Spiro COLORINOX	II a klasė

<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
A004B02103501	Spiro COLORINOX	II a klasė
A004B02104001	Spiro COLORINOX	II a klasė
A004B02105001	Spiro COLORINOX	II a klasė
A004B02106001	Spiro COLORINOX	II a klasė
A004B02190001	Spiro COLORINOX	II a klasė
A004C02102001	Spiro Short (trumpas) Barbed Nervbroach	II a klasė
A004C02102501	Spiro Short (trumpas) Barbed Nervbroach	II a klasė
A004C02103001	Spiro Short (trumpas) Barbed Nervbroach	II a klasė
A004C02103501	Spiro Short (trumpas) Barbed Nervbroach	II a klasė
A004C02104001	Spiro Short (trumpas) Barbed Nervbroach	II a klasė
A004C02105001	Spiro Short (trumpas) Barbed Nervbroach	II a klasė
A004C02106001	Spiro Short (trumpas) Barbed Nervbroach	II a klasė
A005032501000	THERMA-CUT	II a klasė
A005032501200	THERMA-CUT	II a klasė
A005032501400	THERMA-CUT	II a klasė
A005032501200	THERMA-CUT	II a klasė
A005032590000	THERMA-CUT	II a klasė
A005131300500	POST SPACE BUR (gražtas)	II a klasė
A005131300700	POST SPACE BUR (gražtas)	II a klasė
A005131900500	POST SPACE BUR (gražtas)	II a klasė
A005131900700	POST SPACE BUR (gražtas)	II a klasė
A011C02101003	K-Flexoreamer sterilus	II a klasė
A011C02101503	K-Flexoreamer sterilus	II a klasė
A011C02102003	K-Flexoreamer sterilus	II a klasė
A011C02102503	K-Flexoreamer sterilus	II a klasė
A011C02103003	K-Flexoreamer sterilus	II a klasė
A011C02103503	K-Flexoreamer sterilus	II a klasė
A011C02104003	K-Flexoreamer sterilus	II a klasė
A011C02104503	K-Flexoreamer sterilus	II a klasė
A011C02105003	K-Flexoreamer sterilus	II a klasė
A011C02105503	K-Flexoreamer sterilus	II a klasė
A011C02106003	K-Flexoreamer sterilus	II a klasė
A011C02107003	K-Flexoreamer sterilus	II a klasė
A011C02108003	K-Flexoreamer sterilus	II a klasė
A011C02109003	K-Flexoreamer sterilus	II a klasė
A011C02190103	K-Flexoreamer sterilus	II a klasė
A011C02190203	K-Flexoreamer sterilus	II a klasė
A011C02500803	K-Flexoreamer sterilus	II a klasė
A011C02501003	K-Flexoreamer sterilus	II a klasė
A011C02501503	K-Flexoreamer sterilus	II a klasė
A011C02512003	K-Flexoreamer sterilus	II a klasė
A011C02502503	K-Flexoreamer sterilus	II a klasė
A011C02503003	K-Flexoreamer sterilus	II a klasė
A011C02503503	K-Flexoreamer sterilus	II a klasė
A011C02504003	K-Flexoreamer sterilus	II a klasė
A011C02504503	K-Flexoreamer sterilus	II a klasė
A011C02505003	K-Flexoreamer sterilus	II a klasė
A011C02505503	K-Flexoreamer sterilus	II a klasė
A011C02506003	K-Flexoreamer sterilus	II a klasė
A011C02507003	K-Flexoreamer sterilus	II a klasė
A011C02508003	K-Flexoreamer sterilus	II a klasė
A011C02509003	K-Flexoreamer sterilus	II a klasė
A011C02590103	K-Flexoreamer sterilus	II a klasė
A011C02590203	K-Flexoreamer sterilus	II a klasė
A011C03101503	K-Flexoreamer sterilus	II a klasė
A011C03102003	K-Flexoreamer sterilus	II a klasė
A011C03102503	K-Flexoreamer sterilus	II a klasė

<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
A011C03103003	K-Flexoreamer sterilus	II a klasė
A011C03103503	K-Flexoreamer sterilus	II a klasė
A011C03104003	K-Flexoreamer sterilus	II a klasė
A011C03190003	K-Flexoreamer sterilus	II a klasė
A011N21801500	PROFILE.04	II a klasė
A011N21801503	PROFILE STERILUS .04	II a klasė
A011N21802000	PROFILE.04	II a klasė
A011N21802003	PROFILE STERILUS .04	II a klasė
A011N21802500	PROFILE.04	II a klasė
A011N21802503	PROFILE STERILUS .04	II a klasė
A011N21803000	PROFILE.04	II a klasė
A011N21803003	PROFILE STERILUS .04	II a klasė
A011N21803500	PROFILE.04	II a klasė
A011N21803503	PROFILE STERILUS .04	II a klasė
A011N21804000	PROFILE.04	II a klasė
A011N21804003	PROFILE STERILUS .04	II a klasė
A011N21890000	PROFILE.04	II a klasė
A011N21890003	PROFILE STERILUS .04	II a klasė
A011N22101500	PROFILE.04	II a klasė
A011N22101503	PROFILE STERILUS .04	II a klasė
A011N22102000	PROFILE.04	II a klasė
A011N22102003	PROFILE STERILUS .04	II a klasė
A011N22102500	PROFILE.04	II a klasė
A011N22102503	PROFILE STERILUS .04	II a klasė
A011N22103000	PROFILE.04	II a klasė
A011N22103003	PROFILE STERILUS .04	II a klasė
A011N22103500	PROFILE.04	II a klasė
A011N22103503	PROFILE STERILUS .04	II a klasė
A011N22104000	PROFILE.04	II a klasė
A011N22104003	PROFILE STERILUS .04	II a klasė
A011N22104500	PROFILE.04	II a klasė
A011N22104503	PROFILE STERILUS .04	II a klasė
A011N22106000	PROFILE.04	II a klasė
A011N22106003	PROFILE STERILUS .04	II a klasė
A011N22109000	PROFILE.04	II a klasė
A011N22109003	PROFILE STERILUS .04	II a klasė
A011N22190000	PROFILE.04	II a klasė
A011N22190003	PROFILE STERILUS .04	II a klasė
A011N22190700	PROFILE.04	II a klasė
A011N22190703	PROFILE STERILUS .04	II a klasė
A011N25101500	PROFILE.04	II a klasė
A011N25101503	PROFILE STERILUS .04	II a klasė
A011N25102000	PROFILE.04	II a klasė
A011N25102003	PROFILE STERILUS .04	II a klasė
A011N25102500	PROFILE.04	II a klasė
A011N25102503	PROFILE STERILUS .04	II a klasė
A011N25103000	PROFILE.04	II a klasė
A011N25103003	PROFILE STERILUS .04	II a klasė
A011N25103500	PROFILE.04	II a klasė
A011N25103503	PROFILE STERILUS .04	II a klasė
A011N25104000	PROFILE.04	II a klasė
A011N25104003	PROFILE STERILUS .04	II a klasė
A011N25104500	PROFILE.04	II a klasė
A011N25104503	PROFILE STERILUS .04	II a klasė
A011N25106000	PROFILE.04	II a klasė
A011N25106003	PROFILE STERILUS .04	II a klasė
A011N25109000	PROFILE.04	II a klasė

<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
A011N25109003	PROFILE STERILUS .04	II a klasė
A011N25190000	PROFILE.04	II a klasė
A011N25190003	PROFILE STERILUS .04	II a klasė
A011N25190700	PROFILE.04	II a klasė
A011N25190703	PROFILE STERILUS .04	II a klasė
A011N23101500	PROFILE.04	II a klasė
A011N23101503	PROFILE STERILUS .04	II a klasė
A011N23102000	PROFILE.04	II a klasė
A011N23102003	PROFILE STERILUS .04	II a klasė
A011N23102500	PROFILE.04	II a klasė
A011N23102503	PROFILE STERILUS .04	II a klasė
A011N23103000	PROFILE.04	II a klasė
A011N23103003	PROFILE STERILUS .04	II a klasė
A011N23103500	PROFILE.04	II a klasė
A011N23103503	PROFILE STERILUS .04	II a klasė
A011N23104000	PROFILE.04	II a klasė
A011N23104003	PROFILE STERILUS .04	II a klasė
A011N23104500	PROFILE.04	II a klasė
A011N23104503	PROFILE STERILUS .04	II a klasė
A011N23106000	PROFILE.04	II a klasė
A011N23106003	PROFILE STERILUS .04	II a klasė
A011N23109000	PROFILE.04	II a klasė
A011N23109003	PROFILE STERILUS .04	II a klasė
A011N23190000	PROFILE.04	II a klasė
A011N23190003	PROFILE STERILUS .04	II a klasė
A011N23190700	PROFILE.04	II a klasė
A011N23190703	PROFILE STERILUS .04	II a klasė
A012C01801503	K-Flexoreamer sterilus	II a klasė
A012C01802003	K-Flexoreamer sterilus	II a klasė
A012C01802503	K-Flexoreamer sterilus	II a klasė
A012C01803003	K-Flexoreamer sterilus	II a klasė
A012C01803503	K-Flexoreamer sterilus	II a klasė
A012C01804003	K-Flexoreamer sterilus	II a klasė
A012C01809003	K-Flexoreamer sterilus	II a klasė
A012C02101603	K-Flexoreamer sterilus	II a klasė
A012C02101803	K-Flexoreamer sterilus	II a klasė
A012C02101003	K-Flexoreamer sterilus	II a klasė
A012C02101503	K-Flexoreamer sterilus	II a klasė
A012C02102003	K-Flexoreamer sterilus	II a klasė
A012C02102503	K-Flexoreamer sterilus	II a klasė
A012C02103003	K-Flexoreamer sterilus	II a klasė
A012C02103503	K-Flexoreamer sterilus	II a klasė
A012C02104003	K-Flexoreamer sterilus	II a klasė
A012C02104503	K-Flexoreamer sterilus	II a klasė
A012C02105003	K-Flexoreamer sterilus	II a klasė
A012C02105503	K-Flexoreamer sterilus	II a klasė
A012C02106003	K-Flexoreamer sterilus	II a klasė
A012C02107003	K-Flexoreamer sterilus	II a klasė
A012C02108003	K-Flexoreamer sterilus	II a klasė
A012C02109003	K-Flexoreamer sterilus	II a klasė
A012C02110003	K-Flexoreamer sterilus	II a klasė
A012C02111003	K-Flexoreamer sterilus	II a klasė
A012C02112003	K-Flexoreamer sterilus	II a klasė
A012C02113003	K-Flexoreamer sterilus	II a klasė
A012C02114003	K-Flexoreamer sterilus	II a klasė
A012C02190003	K-Flexoreamer sterilus	II a klasė
A012C02190103	K-Flexoreamer sterilus	II a klasė

<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
A012C02190203	K-Flexoreamer sterilus	II a klasė
A012C02501603	K-Flexoreamer sterilus	II a klasė
A012C02501803	K-Flexoreamer sterilus	II a klasė
A012C02501003	K-Flexoreamer sterilus	II a klasė
A012C02501503	K-Flexoreamer sterilus	II a klasė
A012C02502003	K-Flexoreamer sterilus	II a klasė
A012C02502503	K-Flexoreamer sterilus	II a klasė
A012C02503003	K-Flexoreamer sterilus	II a klasė
A012C02503503	K-Flexoreamer sterilus	II a klasė
A012C02504003	K-Flexoreamer sterilus	II a klasė
A012C02504503	K-Flexoreamer sterilus	II a klasė
A012C02505003	K-Flexoreamer sterilus	II a klasė
A012C02505503	K-Flexoreamer sterilus	II a klasė
A012C02506003	K-Flexoreamer sterilus	II a klasė
A012C02507003	K-Flexoreamer sterilus	II a klasė
A012C02508003	K-Flexoreamer sterilus	II a klasė
A012C02509003	K-Flexoreamer sterilus	II a klasė
A012C02510003	K-Flexoreamer sterilus	II a klasė
A012C02511003	K-Flexoreamer sterilus	II a klasė
A012C02512003	K-Flexoreamer sterilus	II a klasė
A012C02513003	K-Flexoreamer sterilus	II a klasė
A012C02514003	K-Flexoreamer sterilus	II a klasė
A012C02590003	K-Flexoreamer sterilus	II a klasė
A012C02590103	K-Flexoreamer sterilus	II a klasė
A012C02590203	K-Flexoreamer sterilus	II a klasė
A012C03101603	K-Flexoreamer sterilus	II a klasė
A012C03101803	K-Flexoreamer sterilus	II a klasė
A012C03101003	K-Flexoreamer sterilus	II a klasė
A012C03101503	K-Flexoreamer sterilus	II a klasė
A012C03102003	K-Flexoreamer sterilus	II a klasė
A012C03102503	K-Flexoreamer sterilus	II a klasė
A012C03103003	K-Flexoreamer sterilus	II a klasė
A012C03103503	K-Flexoreamer sterilus	II a klasė
A012C03104003	K-Flexoreamer sterilus	II a klasė
A012C03104503	K-Flexoreamer sterilus	II a klasė
A012C03105003	K-Flexoreamer sterilus	II a klasė
A012C03105503	K-Flexoreamer sterilus	II a klasė
A012C03106003	K-Flexoreamer sterilus	II a klasė
A012C03107003	K-Flexoreamer sterilus	II a klasė
A012C03108003	K-Flexoreamer sterilus	II a klasė
A012C03109003	K-Flexoreamer sterilus	II a klasė
A012C03110003	K-Flexoreamer sterilus	II a klasė
A012C03111003	K-Flexoreamer sterilus	II a klasė
A012C03112003	K-Flexoreamer sterilus	II a klasė
A012C03113003	K-Flexoreamer sterilus	II a klasė
A012C03114003	K-Flexoreamer sterilus	II a klasė
A012C03190003	K-Flexoreamer sterilus	II a klasė
A012C03190103	K-Flexoreamer sterilus	II a klasė
A012C03190203	K-Flexoreamer sterilus	II a klasė
A012D01800603	K-File COLORINOX sterilus	II a klasė
A012D01800803	K-File COLORINOX sterilus	II a klasė
A012D01801003	K-File COLORINOX sterilus	II a klasė
A012D02100603	K-File COLORINOX sterilus	II a klasė
A012D02100803	K-File COLORINOX sterilus	II a klasė
A012D02101003	K-File COLORINOX sterilus	II a klasė
A012D02101503	K-File COLORINOX sterilus	II a klasė
A012D02102003	K-File COLORINOX sterilus	II a klasė





<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
A012N02101503	K-File NITIFLEX sterilus	II a klasė
A012N02102003	K-File NITIFLEX sterilus	II a klasė
A012N02102503	K-File NITIFLEX sterilus	II a klasė
A012N02103003	K-File NITIFLEX sterilus	II a klasė
A012N02103503	K-File NITIFLEX sterilus	II a klasė
A012N02104003	K-File NITIFLEX sterilus	II a klasė
A012N02104503	K-File NITIFLEX sterilus	II a klasė
A012N02105003	K-File NITIFLEX sterilus	II a klasė
A012N02105503	K-File NITIFLEX sterilus	II a klasė
A012N02106003	K-File NITIFLEX sterilus	II a klasė
A012N02190003	K-File NITIFLEX sterilus	II a klasė
A012N02501503	K-File NITIFLEX sterilus	II a klasė
A012N02502003	K-File NITIFLEX sterilus	II a klasė
A012N02502503	K-File NITIFLEX sterilus	II a klasė
A012N02503003	K-File NITIFLEX sterilus	II a klasė
A012N02503503	K-File NITIFLEX sterilus	II a klasė
A012N02504003	K-File NITIFLEX sterilus	II a klasė
A012N02504503	K-File NITIFLEX sterilus	II a klasė
A012N02505003	K-File NITIFLEX sterilus	II a klasė
A012N02505503	K-File NITIFLEX sterilus	II a klasė
A012N02506003	K-File NITIFLEX sterilus	II a klasė
A012N02590003	K-File NITIFLEX sterilus	II a klasė
A012X01800603	C+ Failai sterilūs kateterizacijai	II a klasė
A012X01800803	C+ Failai sterilūs kateterizacijai	II a klasė
A012X01801003	C+ Failai sterilūs kateterizacijai	II a klasė
A012X01801503	C+ Failai sterilūs kateterizacijai	II a klasė
A012X01802003	C+ Failai sterilūs kateterizacijai	II a klasė
A012X01890003	C+ Failai sterilūs kateterizacijai	II a klasė
A012X02100603	C+ Failai sterilūs kateterizacijai	II a klasė
A012X02100803	C+ Failai sterilūs kateterizacijai	II a klasė
A012X02101003	C+ Failai sterilūs kateterizacijai	II a klasė
A012X02101503	C+ Failai sterilūs kateterizacijai	II a klasė
A012X02190003	C+ Failai sterilūs kateterizacijai	II a klasė
A012X02500603	C+ Failai sterilūs kateterizacijai	II a klasė
A012X02500803	C+ Failai sterilūs kateterizacijai	II a klasė
A012X02501003	C+ Failai sterilūs kateterizacijai	II a klasė
A012X02501503	C+ Failai sterilūs kateterizacijai	II a klasė
A012X02590003	C+ Failai sterilūs kateterizacijai	II a klasė
A016430000100	Endo access grąžtas	II a klasė
A016430000200	Endo access grąžtas	II a klasė
A016430000300	Endo access grąžtas	II a klasė
A016430090000	Endo access grąžtas	II a klasė
A016532100100	Diamendo	II a klasė
A016532500200	Diamendo	II a klasė
A016D02100803	Hedstroem File Colorinox sterilus	II a klasė
A016D02101003	Hedstroem File Colorinox sterilus	II a klasė
A016D02101503	Hedstroem File Colorinox sterilus	II a klasė
A016D02102003	Hedstroem File Colorinox sterilus	II a klasė
A016D02102503	Hedstroem File Colorinox sterilus	II a klasė
A016D02103003	Hedstroem File Colorinox sterilus	II a klasė
A016D02103503	Hedstroem File Colorinox sterilus	II a klasė
A016D02104003	Hedstroem File Colorinox sterilus	II a klasė
A016D02104503	Hedstroem File Colorinox sterilus	II a klasė
A016D02105003	Hedstroem File Colorinox sterilus	II a klasė
A016D02105503	Hedstroem File Colorinox sterilus	II a klasė
A016D02106003	Hedstroem File Colorinox sterilus	II a klasė
A016D02107003	Hedstroem File Colorinox sterilus	II a klasė



<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
A016D03101003	Hedstroem File Colorinox sterilus	II a klasė
A016D03101503	Hedstroem File Colorinox sterilus	II a klasė
A016D03102003	Hedstroem File Colorinox sterilus	II a klasė
A016D03102503	Hedstroem File Colorinox sterilus	II a klasė
A016D03103003	Hedstroem File Colorinox sterilus	II a klasė
A016D03103503	Hedstroem File Colorinox sterilus	II a klasė
A016D03104003	Hedstroem File Colorinox sterilus	II a klasė
A016D03104503	Hedstroem File Colorinox sterilus	II a klasė
A016D03105003	Hedstroem File Colorinox sterilus	II a klasė
A016D03105503	Hedstroem File Colorinox sterilus	II a klasė
A016D03106003	Hedstroem File Colorinox sterilus	II a klasė
A016D03107003	Hedstroem File Colorinox sterilus	II a klasė
A016D03108003	Hedstroem File Colorinox sterilus	II a klasė
A016D03109003	Hedstroem File Colorinox sterilus	II a klasė
A016D03110003	Hedstroem File Colorinox sterilus	II a klasė
A016D03111003	Hedstroem File Colorinox sterilus	II a klasė
A016D03112003	Hedstroem File Colorinox sterilus	II a klasė
A016D03113003	Hedstroem File Colorinox sterilus	II a klasė
A016D03114003	Hedstroem File Colorinox sterilus	II a klasė
A016D03190003	Hedstroem File Colorinox sterilus	II a klasė
A016D03190103	Hedstroem File Colorinox sterilus	II a klasė
A016D03190203	Hedstroem File Colorinox sterilus	II a klasė
A016G02101000	File Hedstroem	II a klasė
A016G02101500	File Hedstroem	II a klasė
A016G02102000	File Hedstroem	II a klasė
A016G02102500	File Hedstroem	II a klasė
A016G02103000	File Hedstroem	II a klasė
A016G02103500	File Hedstroem	II a klasė
A016G02104000	File Hedstroem	II a klasė
A016G02104500	File Hedstroem	II a klasė
A016G02105000	File Hedstroem	II a klasė
A016G02105500	File Hedstroem	II a klasė
A016G02108000	File Hedstroem	II a klasė
A016G02190000	File Hedstroem	II a klasė
A016G02501000	File Hedstroem	II a klasė
A016G02501500	File Hedstroem	II a klasė
A016G02502000	File Hedstroem	II a klasė
A016G02502500	File Hedstroem	II a klasė
A016G02503000	File Hedstroem	II a klasė
A016G02503500	File Hedstroem	II a klasė
A016G02504000	File Hedstroem	II a klasė
A016G02504500	File Hedstroem	II a klasė
A016G02505000	File Hedstroem	II a klasė
A016G02505500	File Hedstroem	II a klasė
A016G02506000	File Hedstroem	II a klasė
A016G02507000	File Hedstroem	II a klasė
A016G02508000	File Hedstroem	II a klasė
A016G02590000	File Hedstroem	II a klasė
A016G02590100	File Hedstroem	II a klasė
A016G02801000	File Hedstroem	II a klasė
A016G02801500	File Hedstroem	II a klasė
A016G02802000	File Hedstroem	II a klasė
A016G02802500	File Hedstroem	II a klasė
A016G02803500	File Hedstroem	II a klasė
A016G02804000	File Hedstroem	II a klasė
A017502502000	Verifier (Tikrintojas)	II a klasė
A017502502003	Verifier (Tikrintojas) sterilus	II a klasė

<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
A017502502500	Verifier (Tikrintojas)	II a klasė
A017502502503	Verifier (Tikrintojas) sterilus	II a klasė
A017502503000	Verifier (Tikrintojas)	II a klasė
A017502503003	Verifier (Tikrintojas) sterilus	II a klasė
A017502503500	Verifier (Tikrintojas)	II a klasė
A017502503503	Verifier (Tikrintojas) sterilus	II a klasė
A017502504000	Verifier (Tikrintojas)	II a klasė
A017502504003	Verifier (Tikrintojas) sterilus	II a klasė
A017502504500	Verifier (Tikrintojas)	II a klasė
A017502504503	Verifier (Tikrintojas) sterilus	II a klasė
A017502505000	Verifier (Tikrintojas)	II a klasė
A017502505003	Verifier (Tikrintojas) sterilus	II a klasė
A017502505500	Verifier (Tikrintojas)	II a klasė
A017502505503	Verifier (Tikrintojas) sterilus	II a klasė
A017502506000	Verifier (Tikrintojas)	II a klasė
A017502506003	Verifier (Tikrintojas) sterilus	II a klasė
A017502507000	Verifier (Tikrintojas)	II a klasė
A017502507003	Verifier (Tikrintojas) sterilus	II a klasė
A017502508000	Verifier (Tikrintojas)	II a klasė
A017502508003	Verifier (Tikrintojas) sterilus	II a klasė
A017502509000	Verifier (Tikrintojas)	II a klasė
A017502509003	Verifier (Tikrintojas) sterilus	II a klasė
A017502590000	Verifier (Tikrintojas)	II a klasė
A017502590003	Verifier (Tikrintojas) sterilus	II a klasė
A017502590100	Verifier (Tikrintojas)	II a klasė
A017502590103	Verifier (Tikrintojas) sterilus	II a klasė
A018202100103	TAPERED FINGER SPREADER STERILUS	II a klasė
A018202100203	TAPERED FINGER SPREADER STERILUS	II a klasė
A018202100303	TAPERED FINGER SPREADER STERILUS	II a klasė
A018202100403	TAPERED FINGER SPREADER STERILUS	II a klasė
A018202190003	TAPERED FINGER SPREADER STERILUS	II a klasė
A018202500103	TAPERED FINGER SPREADER STERILUS	II a klasė
A018202500203	TAPERED FINGER SPREADER STERILUS	II a klasė
A018202500303	TAPERED FINGER SPREADER STERILUS	II a klasė
A018202500403	TAPERED FINGER SPREADER STERILUS	II a klasė
A018202590003	TAPERED FINGER SPREADER STERILUS	II a klasė
A022S21500300	Lentulo trumpas, ilgas ir x-ilgas	II a klasė
A022S21500303	Lentulo sterilus Pastos nešėjas	II a klasė
A022S21700100	Lentulo trumpas, ilgas ir x-ilgas	II a klasė
A022S21700101	Lentulo sterilus Pastos nešėjas	II a klasė
A022S21700103	Lentulo sterilus Pastos nešėjas	II a klasė
A022S21700200	Lentulo trumpas, ilgas ir x-ilgas	II a klasė
A022S21700203	Lentulo sterilus Pastos nešėjas	II a klasė
A022S21700300	Lentulo trumpas, ilgas ir x-ilgas	II a klasė
A022S21700301	Lentulo sterilus Pastos nešėjas	II a klasė
A022S21700303	Lentulo sterilus Pastos nešėjas	II a klasė
A022S21700400	Lentulo trumpas, ilgas ir x-ilgas	II a klasė
A022S21700401	Lentulo sterilus Pastos nešėjas	II a klasė
A022S21700403	Lentulo sterilus Pastos nešėjas	II a klasė
A022S22100100	Lentulo trumpas, ilgas ir x-ilgas	II a klasė
A022S22100101	Lentulo sterilus Pastos nešėjas	II a klasė
A022S22100103	Lentulo sterilus Pastos nešėjas	II a klasė
A022S22100200	Lentulo trumpas, ilgas ir x-ilgas	II a klasė
A022S22100201	Lentulo trumpas, ilgas ir x-ilgas	II a klasė
A022S22100203	Lentulo sterilus Pastos nešėjas	II a klasė

<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
A022S22100300	Lentulo trumpas, ilgas ir x-ilgas	II a klasė
A022S22100301	Lentulo sterilus Pastos nešėjas	II a klasė
A022S22100303	Lentulo sterilus Pastos nešėjas	II a klasė
A022S22100400	Lentulo trumpas, ilgas ir x-ilgas	II a klasė
A022S22100401	Lentulo sterilus Pastos nešėjas	II a klasė
A022S22100403	Lentulo sterilus Pastos nešėjas	II a klasė
A022S22500100	Lentulo trumpas, ilgas ir x-ilgas	II a klasė
A022S22500101	Lentulo sterilus Pastos nešėjas	II a klasė
A022S22500103	Lentulo sterilus Pastos nešėjas	II a klasė
A022S22500200	Lentulo trumpas, ilgas ir x-ilgas	II a klasė
A022S22500201	Lentulo trumpas, ilgas ir x-ilgas	II a klasė
A022S22500203	Lentulo sterilus Pastos nešėjas	II a klasė
A022S22500300	Lentulo trumpas, ilgas ir x-ilgas	II a klasė
A022S22500301	Lentulo sterilus Pastos nešėjas	II a klasė
A022S22500303	Lentulo sterilus Pastos nešėjas	II a klasė
A022S22500400	Lentulo trumpas, ilgas ir x-ilgas	II a klasė
A022S22500401	Lentulo sterilus Pastos nešėjas	II a klasė
A022S22500403	Lentulo sterilus Pastos nešėjas	II a klasė
A024202102500	GUTTA-KONDENSATORIUS	II a klasė
A024202102503	GUTTA-KONDENSATORIUS STERILUS	II a klasė
A024202103000	GUTTA-KONDENSATORIUS	II a klasė
A024202103003	GUTTA-KONDENSATORIUS STERILUS	II a klasė
A024202103500	GUTTA-KONDENSATORIUS	II a klasė
A024202103503	GUTTA-KONDENSATORIUS STERILUS	II a klasė
A024202104000	GUTTA-KONDENSATORIUS	II a klasė
A024202104003	GUTTA-KONDENSATORIUS STERILUS	II a klasė
A024202104500	GUTTA-KONDENSATORIUS	II a klasė
A024202104503	GUTTA-KONDENSATORIUS STERILUS	II a klasė
A024202105000	GUTTA-KONDENSATORIUS	II a klasė
A024202105003	GUTTA-KONDENSATORIUS STERILUS	II a klasė
A024202105500	GUTTA-KONDENSATORIUS	II a klasė
A024202105503	GUTTA-KONDENSATORIUS STERILUS	II a klasė
A024202106000	GUTTA-KONDENSATORIUS	II a klasė
A024202106003	GUTTA-KONDENSATORIUS STERILUS	II a klasė
A024202107000	GUTTA-KONDENSATORIUS	II a klasė
A024202107003	GUTTA-KONDENSATORIUS STERILUS	II a klasė
A024202108000	GUTTA-KONDENSATORIUS	II a klasė
A024202108003	GUTTA-KONDENSATORIUS STERILUS	II a klasė
A024202109000	GUTTA-KONDENSATORIUS	II a klasė
A024202502500	GUTTA-KONDENSATORIUS	II a klasė
A024202502503	GUTTA-KONDENSATORIUS STERILUS	II a klasė
A024202502500	GUTTA-KONDENSATORIUS	II a klasė
A024205503003	GUTTA-KONDENSATORIUS STERILUS	II a klasė
A024202503500	GUTTA-KONDENSATORIUS	II a klasė
A024202503503	GUTTA-KONDENSATORIUS STERILUS	II a klasė
A024202504000	GUTTA-KONDENSATORIUS	II a klasė
A024202504003	GUTTA-KONDENSATORIUS STERILUS	II a klasė
A024202504500	GUTTA-KONDENSATORIUS	II a klasė
A024202504503	GUTTA-KONDENSATORIUS STERILUS	II a klasė
A024202505000	GUTTA-KONDENSATORIUS	II a klasė
A024202505003	GUTTA-KONDENSATORIUS STERILUS	II a klasė
A024202505500	GUTTA-KONDENSATORIUS	II a klasė
A024202505503	GUTTA-KONDENSATORIUS STERILUS	II a klasė
A024202506000	GUTTA-KONDENSATORIUS	II a klasė
A024202506003	GUTTA-KONDENSATORIUS STERILUS	II a klasė

<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
A024202507000	GUTTA-KONDENSATORIUS	II a klasė
A024202507003	GUTTA-KONDENSATORIUS STERILUS	II a klasė
A024202508000	GUTTA-KONDENSATORIUS	II a klasė
A024202508003	GUTTA-KONDENSATORIUS STERILUS	II a klasė
A024202509000	GUTTA-KONDENSATORIUS	II a klasė
A024202190000	GUTTA-KONDENSATORIŲ rinkinys	II a klasė
A024202590000	GUTTA-KONDENSATORIŲ rinkinys	II a klasė
A027400000000	Flexokit Full	II a klasė
A030500000000	Cavity Access rinkinys	II a klasė
A030500000100	Cavity Access rinkinys	II a klasė
A034422101500	ProFile .02	II a klasė
A034422101503	ProFile STERILUS .02	II a klasė
A034422102000	ProFile .02	II a klasė
A034422102003	ProFile STERILUS .02	II a klasė
A034422102500	ProFile .02	II a klasė
A034422102503	ProFile STERILUS .02	II a klasė
A034422103000	ProFile .02	II a klasė
A034422103003	ProFile STERILUS .02	II a klasė
A034422103500	ProFile .02	II a klasė
A034422103503	ProFile STERILUS .02	II a klasė
A034422104000	ProFile .02	II a klasė
A034422104003	ProFile STERILUS .02	II a klasė
A034422190000	ProFile .02	II a klasė
A034422190003	ProFile STERILUS .02	II a klasė
A034422501500	ProFile .02	II a klasė
A034422501503	ProFile STERILUS .02	II a klasė
A034422502000	ProFile .02	II a klasė
A034422502003	ProFile STERILUS .02	II a klasė
A034422502500	ProFile .02	II a klasė
A034422502503	ProFile STERILUS .02	II a klasė
A034422503000	ProFile .02	II a klasė
A034422503003	ProFile STERILUS .02	II a klasė
A034422503500	ProFile .02	II a klasė
A034422503503	ProFile STERILUS .02	II a klasė
A034422504000	ProFile .02	II a klasė
A034422504003	ProFile STERILUS .02	II a klasė
A034422590000	ProFile .02	II a klasė
A034422590003	ProFile STERILUS .02	II a klasė
A034521801500	ProFile .06	II a klasė
A034521801503	ProFile STERILUS .06	II a klasė
A034521802000	ProFile .06	II a klasė
A034521802003	ProFile STERILUS .06	II a klasė
A034521802500	ProFile .06	II a klasė
A034521802503	ProFile STERILUS .06	II a klasė
A034521803000	ProFile .06	II a klasė
A034521803003	ProFile STERILUS .06	II a klasė
A034521803500	ProFile .06	II a klasė
A034521803503	ProFile STERILUS .06	II a klasė
A034521804000	ProFile .06	II a klasė
A034521804003	ProFile STERILUS .06	II a klasė
A034521890000	ProFile .06	II a klasė
A034521890003	ProFile STERILUS .06	II a klasė
A034522101500	ProFile .06	II a klasė
A034522101503	ProFile STERILUS .06	II a klasė
A034522102000	ProFile .06	II a klasė
A034522102003	ProFile STERILUS .06	II a klasė
A034522102500	ProFile .06	II a klasė

<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
A034522102503	ProFile STERILUS .06	II a klasė
A034522103000	ProFile .06	II a klasė
A034522103003	ProFile STERILUS .06	II a klasė
A03452213500	ProFile .06	II a klasė
A034522103503	ProFile STERILUS .06	II a klasė
A034522104000	ProFile .06	II a klasė
A034522104003	ProFile STERILUS .06	II a klasė
A034522190000	ProFile .06	II a klasė
A034522190003	ProFile STERILUS .06	II a klasė
A034522501500	ProFile .06	II a klasė
A034522501503	ProFile STERILUS .06	II a klasė
A034522502000	ProFile .06	II a klasė
A034522502003	ProFile STERILUS .06	II a klasė
A034522502500	ProFile .06	II a klasė
A034522502503	ProFile STERILUS .06	II a klasė
A034522503000	ProFile .06	II a klasė
A034522503003	ProFile STERILUS .06	II a klasė
A03452253500	ProFile .06	II a klasė
A034522503503	ProFile STERILUS .06	II a klasė
A034522504000	ProFile .06	II a klasė
A034522504003	ProFile STERILUS .06	II a klasė
A034522590000	ProFile .06	II a klasė
A034522590003	ProFile STERILUS .06	II a klasė
A035802590000	ProFile O.S. + .06 + .04	II a klasė
A035921900100	ProFile O.S.	II a klasė
A035921900103	ProFile STERILUS O.S.	II a klasė
A035921900200	ProFile O.S.	II a klasė
A035921900203	ProFile STERILUS O.S.	II a klasė
A035921900300	ProFile O.S.	II a klasė
A035921900303	ProFile STERILUS O.S.	II a klasė
A035921900400	ProFile O.S.	II a klasė
A035921900403	ProFile STERILUS O.S.	II a klasė
A035921900500	ProFile O.S.	II a klasė
A035921900503	ProFile STERILUS O.S.	II a klasė
A035921900600	ProFile O.S.	II a klasė
A035921900603	ProFile STERILUS O.S.	II a klasė
A035921990000	ProFile O.S.	II a klasė
A035921990003	ProFile STERILUS O.S.	II a klasė
A036102101000	Unifile	II a klasė
A036102101500	Unifile	II a klasė
A036102102000	Unifile	II a klasė
A036102102500	Unifile	II a klasė
A036102103000	Unifile	II a klasė
A036102103500	Unifile	II a klasė
A036102105000	Unifile	II a klasė
A036102105500	Unifile	II a klasė
A036102106000	Unifile	II a klasė
A036102107000	Unifile	II a klasė
A036102190000	Unifile	II a klasė
A036102190100	Unifile	II a klasė
A036102191000	Unifile	II a klasė
A036102501000	Unifile	II a klasė
A036102501500	Unifile	II a klasė
A036102502000	Unifile	II a klasė
A036102503000	Unifile	II a klasė
A036102504000	Unifile	II a klasė
A036102504500	Unifile	II a klasė

<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
A036102505000	Unifile	II a klasė
A036102505500	Unifile	II a klasė
A036102506000	Unifile	II a klasė
A036102507000	Unifile	II a klasė
A036102508000	Unifile	II a klasė
A036102590000	Unifile	II a klasė
A036102590100	Unifile	II a klasė
A036102591000	Unifile	II a klasė
A036200000000	PROFILE Organizer	II a klasė
A036200000900	PROFILE Organizer	II a klasė
A036302100100	PROFILE	II a klasė
A036302500200	PROFILE	II a klasė
A040922100100	PROTAPER UNIVERSAL SEQ BASE	II a klasė
A040922100103	PROTAPER UNIVERSAL Sterilus rinkinys SEQ BASE	II a klasė
A040922190100	PROTAPER UNIVERSAL SEQ BASE	II a klasė
A040922190103	PROTAPER UNIVERSAL Sterilus rinkinys SEQ BASE	II a klasė
A040922500200	PROTAPER UNIVERSAL SEQ BASE	II a klasė
A040922500203	PROTAPER UNIVERSAL Sterilus rinkinys SEQ BASE	II a klasė
A040922590100	PROTAPER UNIVERSAL SEQ BASE	II a klasė
A040922590103	PROTAPER UNIVERSAL Sterilus rinkinys SEQ BASE	II a klasė
A040923100300	PROTAPER UNIVERSAL SEQ BASE	II a klasė
A040923100303	PROTAPER UNIVERSAL Sterilus rinkinys SEQ BASE	II a klasė
A040923190100	PROTAPER UNIVERSAL SEQ BASE	II a klasė
A040923190103	PROTAPER UNIVERSAL Sterilus rinkinys SEQ BASE	II a klasė
A041021910100	PROTAPER UNIVERSAL Shaping failai	II a klasė
A041021910103	PROTAPER UNIVERSAL Shaping failai sterilūs	II a klasė
A041022110100	PROTAPER UNIVERSAL Shaping failai	II a klasė
A041022110103	PROTAPER UNIVERSAL Shaping failai sterilūs	II a klasė
A041022110200	PROTAPER UNIVERSAL Shaping failai	II a klasė
A041022110203	PROTAPER UNIVERSAL Shaping failai sterilūs	II a klasė
A041022510100	PROTAPER UNIVERSAL Shaping failai	II a klasė
A041022510103	PROTAPER UNIVERSAL Shaping failai sterilūs	II a klasė
A041022510200	PROTAPER UNIVERSAL Shaping failai	II a klasė
A041022510203	PROTAPER UNIVERSAL Shaping failai sterilūs	II a klasė
A041023110100	PROTAPER UNIVERSAL Shaping failai	II a klasė
A041023110103	PROTAPER UNIVERSAL Shaping failai sterilūs	II a klasė
A041023110200	PROTAPER UNIVERSAL Shaping failai	II a klasė
A041023110203	PROTAPER UNIVERSAL Shaping failai sterilūs	II a klasė
A041122110100	PROTAPER UNIVERSAL Finishing failai	II a klasė
A041122110103	PROTAPER UNIVERSAL Finishing failai sterilūs	II a klasė
A041122110200	PROTAPER UNIVERSAL Finishing failai	II a klasė
A041122110203	PROTAPER UNIVERSAL Finishing failai sterilūs	II a klasė
A041122110300	PROTAPER UNIVERSAL Finishing failai	II a klasė
A041122110303	PROTAPER UNIVERSAL Finishing failai sterilūs	II a klasė
A041122110400	PROTAPER UNIVERSAL Finishing failai	II a klasė
A041122110403	PROTAPER UNIVERSAL Finishing failai sterilūs	II a klasė
A041122110500	PROTAPER UNIVERSAL Finishing failai	II a klasė
A041122110503	PROTAPER UNIVERSAL Finishing failai sterilūs	II a klasė
A041122510100	PROTAPER UNIVERSAL Finishing failai	II a klasė
A041122510103	PROTAPER UNIVERSAL Finishing failai sterilūs	II a klasė
A041122510200	PROTAPER UNIVERSAL Finishing failai	II a klasė
A041122510203	PROTAPER UNIVERSAL Finishing failai sterilūs	II a klasė
A041122510300	PROTAPER UNIVERSAL Finishing failai	II a klasė
A041122510303	PROTAPER UNIVERSAL Finishing failai sterilūs	II a klasė
A041122510400	PROTAPER UNIVERSAL Finishing failai	II a klasė
A041122510403	PROTAPER UNIVERSAL Finishing failai sterilūs	II a klasė
A041122510500	PROTAPER UNIVERSAL Finishing failai	II a klasė

Prod.nr.	Produkto žymėjimas	Produkto klasė
A041122510503	PROTAPER UNIVERSAL Finishing failai sterilūs	II a klasė
A041123110100	PROTAPER UNIVERSAL Finishing failai	II a klasė
A041123110103	PROTAPER UNIVERSAL Finishing failai sterilūs	II a klasė
A041123110200	PROTAPER UNIVERSAL Finishing failai	II a klasė
A041123110203	PROTAPER UNIVERSAL Finishing failai sterilūs	II a klasė
A041123110300	PROTAPER UNIVERSAL Finishing failai	II a klasė
A041123110303	PROTAPER UNIVERSAL Finishing failai sterilūs	II a klasė
A041123110400	PROTAPER UNIVERSAL Finishing failai	II a klasė
A041123110403	PROTAPER UNIVERSAL Finishing failai sterilūs	II a klasė
A041123110500	PROTAPER UNIVERSAL Finishing failai	II a klasė
A041123110503	PROTAPER UNIVERSAL Finishing failai sterilūs	II a klasė
A041522190100	Pradinis rinkinys PROTAPER UNIVERSAL	II a klasė
A041522590100	Pradinis rinkinys PROTAPER UNIVERSAL	II a klasė
A041523190100	Pradinis rinkinys PROTAPER UNIVERSAL	II a klasė
A066000000000	START-X®	II a klasė
A0660000000100	START-X®	II a klasė
A0660000000200	START-X®	II a klasė
A0660000000300	START-X®	II a klasė
A0660000000400	START-X®	II a klasė
A0660000000500	START-X®	II a klasė
A066000090000	START-X®	II a klasė
A066100000100	START-X®	II a klasė
A066100000200	START-X®	II a klasė
A066100000300	START-X®	II a klasė
A066100000400	START-X®	II a klasė
A066100000500	START-X®	II a klasė
A066100090000	START-X®	II a klasė
A070022100L01	WAVEONE reciprokiniai failai sterilūs	II a klasė
A070022100P01	WAVEONE reciprokiniai failai sterilūs	II a klasė
A070022100S01	WAVEONE reciprokiniai failai sterilūs	II a klasė
A070022190A01	WAVEONE Rinkinys reciprokinių failų sterilūs	II a klasė
A070022500L01	WAVEONE reciprokiniai failai sterilūs	II a klasė
A070022500P01	WAVEONE reciprokiniai failai sterilūs	II a klasė
A070022500S01	WAVEONE reciprokiniai failai sterilūs	II a klasė
A070022590A01	WAVEONE Rinkinys reciprokinių failų sterilūs	II a klasė
A070023100L01	WAVEONE reciprokiniai failai sterilūs	II a klasė
A070023100P01	WAVEONE reciprokiniai failai sterilūs	II a klasė
A070023100S01	WAVEONE reciprokiniai failai sterilūs	II a klasė
A070023190A01	WAVEONE Rinkinys reciprokinių failų sterilūs	II a klasė
A070622100L01	WAVEONE reciprokiniai failai sterilūs	II a klasė
A070622100P01	WAVEONE reciprokiniai failai sterilūs	II a klasė
A070622100S01	WAVEONE reciprokiniai failai sterilūs	II a klasė
A070622190A01	WAVEONE Rinkinys reciprokinių failų sterilūs	II a klasė
A070622500L01	WAVEONE reciprokiniai failai sterilūs	II a klasė
A070622500P01	WAVEONE reciprokiniai failai sterilūs	II a klasė
A070622500S01	WAVEONE reciprokiniai failai sterilūs	II a klasė
A070622590A01	WAVEONE Rinkinys reciprokinių failų sterilūs	II a klasė
A070623100L01	WAVEONE reciprokiniai failai sterilūs	II a klasė
A070623100P01	WAVEONE reciprokiniai failai sterilūs	II a klasė
A070623100S01	WAVEONE reciprokiniai failai sterilūs	II a klasė
A070623190A01	WAVEONE Rinkinys reciprokinių failų sterilūs	II a klasė
A0751225G90D03	WAVEONE GOLD One reciprokiniai failai sterilūs	II a klasė
A0753221GOL03	WAVEONE GOLD reciprokiniai failai sterilūs	II a klasė
A0753221GOM03	WAVEONE GOLD reciprokiniai failai sterilūs	II a klasė
A0753221GOP03	WAVEONE GOLD reciprokiniai failai sterilūs	II a klasė

Prod.nr.	Produkto žymėjimas	Produkto klasė
A0753221GOS03	WAVEONE GOLD reciprokiniai failai sterilūs	II a klasė
A0753225GOL03	WAVEONE GOLD reciprokiniai failai sterilūs	II a klasė
A0753225GOM03	WAVEONE GOLD reciprokiniai failai sterilūs	II a klasė
A0753225GOP03	WAVEONE GOLD reciprokiniai failai sterilūs	II a klasė
A0753225GOS03	WAVEONE GOLD reciprokiniai failai sterilūs	II a klasė
A0753231GOL03	WAVEONE GOLD reciprokiniai failai sterilūs	II a klasė
A0753231GOM03	WAVEONE GOLD reciprokiniai failai sterilūs	II a klasė
A0753231GOP03	WAVEONE GOLD reciprokiniai failai sterilūs	II a klasė
A0753231GOS03	WAVEONE GOLD reciprokiniai failai sterilūs	II a klasė
A0754221G9A03	WAVEONE GOLD Rinkinys reciprokinių failų sterilūs	II a klasė
A0754225G9A03	WAVEONE GOLD Rinkinys reciprokinių failų sterilūs	II a klasė
A0754231G9A03	WAVEONE GOLD Rinkinys reciprokinių failų sterilūs	II a klasė
A0756221GOL03	WAVEONE GOLD reciprokiniai failai sterilūs	II a klasė
A0756221GOM03	WAVEONE GOLD reciprokiniai failai sterilūs	II a klasė
A0756221GOP03	WAVEONE GOLD reciprokiniai failai sterilūs	II a klasė
A0756221GOS03	WAVEONE GOLD reciprokiniai failai sterilūs	II a klasė
A0756225GOL03	WAVEONE GOLD reciprokiniai failai sterilūs	II a klasė
A0756225GOM03	WAVEONE GOLD reciprokiniai failai sterilūs	II a klasė
A0756225GOP03	WAVEONE GOLD reciprokiniai failai sterilūs	II a klasė
A0756225GOS03	WAVEONE GOLD reciprokiniai failai sterilūs	II a klasė
A0756231GOL03	WAVEONE GOLD reciprokiniai failai sterilūs	II a klasė
A0756231GOM03	WAVEONE GOLD reciprokiniai failai sterilūs	II a klasė
A0756231GOP03	WAVEONE GOLD reciprokiniai failai sterilūs	II a klasė
A0756231GOS03	WAVEONE GOLD reciprokiniai failai sterilūs	II a klasė
A080021900A03	PROTAPER NEXT XA	II a klasė
A080022100103	PROTAPER NEXT	II a klasė
A080022100203	PROTAPER NEXT	II a klasė
A080022100303	PROTAPER NEXT	II a klasė
A080022100403	PROTAPER NEXT	II a klasė
A080022100503	PROTAPER NEXT	II a klasė
A08002219A003	PROTAPER NEXT rinkinys	II a klasė
A080022500103	PROTAPER NEXT	II a klasė
A080022500203	PROTAPER NEXT	II a klasė
A080022500303	PROTAPER NEXT	II a klasė
A080022500403	PROTAPER NEXT	II a klasė
A080022500503	PROTAPER NEXT	II a klasė
A08002259A003	PROTAPER NEXT rinkinys	II a klasė
A080023100103	PROTAPER NEXT	II a klasė
A080023100203	PROTAPER NEXT	II a klasė
A080023100303	PROTAPER NEXT	II a klasė
A080023100403	PROTAPER NEXT	II a klasė
A080023100503	PROTAPER NEXT	II a klasė
A08002319A003	PROTAPER NEXT rinkinys	II a klasė
A08022259D003	PROTAPER NEXT Demo	II a klasė
A080321900A03	PROTAPER NEXT XA	II a klasė
A080322100103	PROTAPER NEXT	II a klasė
A080322100203	PROTAPER NEXT	II a klasė
A080322100303	PROTAPER NEXT	II a klasė
A080322100403	PROTAPER NEXT	II a klasė
A080322100503	PROTAPER NEXT	II a klasė
A08032219A003	PROTAPER NEXT rinkinys	II a klasė
A080322500103	PROTAPER NEXT	II a klasė
A080322500203	PROTAPER NEXT	II a klasė
A080322500303	PROTAPER NEXT	II a klasė
A080322500403	PROTAPER NEXT	II a klasė
A080322500503	PROTAPER NEXT	II a klasė
A08032259A003	PROTAPER NEXT rinkinys	II a klasė

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A080323100103	PROTAPER NEXT	II a klasė
A080323100203	PROTAPER NEXT	II a klasė
A080323100303	PROTAPER NEXT	II a klasė
A080323100403	PROTAPER NEXT	II a klasė
A080323100503	PROTAPER NEXT	II a klasė
A08032319A003	PROTAPER NEXT rinkinys	II a klasė
A082100023100	PRORINSE	II a klasė
A082100023400	PRORINSE	II a klasė
A082100024100	PRORINSE	II a klasė
A082100024400	PRORINSE	II a klasė
A082100025100	PRORINSE	II a klasė
A082100025400	PRORINSE	II a klasė
A082100028100	PRORINSE	II a klasė
A082100028400	PRORINSE	II a klasė
A082100030100	PRORINSE	II a klasė
A082100030400	PRORINSE	II a klasė
A092322100103	PROGLIDER	II a klasė
A092322500103	PROGLIDER	II a klasė
A092323100103	PROGLIDER	II a klasė
A09242259D003	PROGLIDER	II a klasė
A092622100103	PROGLIDER	II a klasė
A092622500103	PROGLIDER	II a klasė
A092623100103	PROGLIDER	II a klasė
A100400000000	X-SMART	II a klasė
A100400000022	X-SMART	II a klasė
A100400000033	X-SMART	II a klasė
A100400000122	X-SMART	II a klasė
A100400000126	X-SMART	II a klasė
A100400000133	X-SMART	II a klasė
A101800000000	WAVEONE endo motoras	II a klasė
A101800000200	WAVEONE endo motoras	II a klasė
A101800000400	WAVEONE endo motoras	II a klasė
A102400000033	X-SMART DUAL	II a klasė
A102800000026	PROPEX II	II a klasė
A102800000033	PROPEX II	II a klasė
A103000000000	PROPEX PIXI™	II a klasė
A103000000100	PROPEX PIXI™	II a klasė
A141021600100	ProTaper Universal Retreatment	II a klasė
A141021600103	ProTaper Universal Retreatment sterilus	II a klasė
A141021800100	ProTaper Universal Retreatment	II a klasė
A141021800103	ProTaper Universal Retreatment sterilus	II a klasė
A141022200100	ProTaper Universal Retreatment	II a klasė
A141022200103	ProTaper Universal Retreatment sterilus	II a klasė
A141220090000	ProTaper Universal Retreatment	II a klasė
A141220090003	ProTaper Universal Retreatment	II a klasė
A141500090000	ProTaper Universal pradinis rinkinys	II a klasė
31050701A0000	X-SMART IQ HANDPIECE (ATGALIO) MOTORAS	II a klasė
A105100000000	X-SMART IQ KAMPINIS ANTGALIS	II a klasė
A230A0000200	Endo rink.univ. Fernando Pessoa	II a klasė
A230A0000300	Endo rink.universidad catolica portuguesa polo de viseu	II a klasė
A230A0000600	Endo rink.universidad del Pais vasco	II a klasė
A230A0000700	Endo rink.universidad de Zaragoza	II a klasė
A230A0001400	Endo rink.ISCS-EGAS Moniz	II a klasė
A230A0001500	Endo rink.universidad San Pablo CEU de Madrid	II a klasė
A230A0002000	Endo rink.universidad Europe Miguel de Cervantes	II a klasė
A305Z0000000	Cavity Access Z rinkinys	II a klasė
A305Z0000100	Cavity Access Z rinkinys	II a klasė

<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
A966000090100	START-X® Band.rinkinys tinkantis Satelec	II a klasė
A966100090100	START-X® Band.rinkinys tinkantis EMS	II a klasė
C005320000100	MOOSER SISTEMA	II a klasė
C005320000200	MOOSER SISTEMA	II a klasė
C005420000100	MOOSER SISTEMA	II a klasė
C005420000200	MOOSER SISTEMA	II a klasė
C005820000000	STP ATKURIAMIEJI KAIŠČIAI (PINS)	II a klasė
C009800000100	RS-RADIX-STIFTE INTEGRAL RINKINYS	II a klasė
C009800000200	RS-RADIX-STIFTE INTEGRAL RINKINYS	II a klasė
C009800000300	RS-RADIX-STIFTE INTEGRAL RINKINYS	II a klasė
C009900000000	RS-RADIX-STIFTE RINKINYS	II a klasė
C010000000100	RS RADIX	II a klasė
C010000000200	RS RADIX	II a klasė
C010000000300	RS RADIX	II a klasė
C010120000100	RS RADIX	II a klasė
C010120000200	RS RADIX	II a klasė
C010120000300	RS RADIX	II a klasė
C010220000100	RS RADIX	II a klasė
C010220000100	RS RADIX	II a klasė
C010220000100	RS RADIX	II a klasė
C019300000100	RADIX-ANKER-STANDART	II a klasė
C019300000200	RADIX-ANKER-STANDART	II a klasė
C019300000300	RADIX-ANKER-STANDART	II a klasė
C020000000000	STP ATKURIAMIEJI KAIŠČIAI (PINS)	II a klasė
C020200800000	MOOSER SISTEMOS INTEGRAL RINKINYS	II a klasė
C020200800100	MOOSER SISTEMOS INTEGRAL RINKINYS	II a klasė
C020201000000	MOOSER SISTEMOS INTEGRAL RINKINYS	II a klasė
C020201000100	MOOSER SISTEMOS INTEGRAL RINKINYS	II a klasė
C020820000000	STP ATKURIAMIEJI KAIŠČIAI (PINS)	II a klasė
C020820000S00	STP ATKURIAMIEJI KAIŠČIAI (PINS)	II a klasė
C021000000000	STP ATKURIAMŲJŲ KAIŠČIŲ (PINS) INTEGRAL RINKINYS	II a klasė
C021100000000	STP PAPILDYMAS	II a klasė
C021220000800	UNIMETRIC X mm	II a klasė
C021220010800	UNIMETRIC X mm	II a klasė
C021220011000	UNIMETRIC X mm	II a klasė
C021220020800	UNIMETRIC X mm	II a klasė
C021220021000	UNIMETRIC X mm	II a klasė
C021220030800	UNIMETRIC X mm	II a klasė
C021220031000	UNIMETRIC X mm	II a klasė
C021320000800	UNIMETRIC X mm	II a klasė
C021320010800	UNIMETRIC X mm	II a klasė
C021320011000	UNIMETRIC X mm	II a klasė
C021320020800	UNIMETRIC X mm	II a klasė
C021320021000	UNIMETRIC X mm	II a klasė
C021320030800	UNIMETRIC X mm	II a klasė
C021320031000	UNIMETRIC X mm	II a klasė
C021420000000	UNIMETRIC	II a klasė
C021620000100	RADIX-ANKER-STANDART	II a klasė
C021620000200	RADIX-ANKER-STANDART	II a klasė
C021620000300	RADIX-ANKER-STANDART	II a klasė
C021620003B00	RADIX-ANKER-STANDART	II a klasė
C021720000100	RADIX-ANKER-STANDART	II a klasė
C021720000200	RADIX-ANKER-STANDART	II a klasė
C021720000300	RADIX-ANKER-STANDART	II a klasė
C021720003B00	RADIX-ANKER-STANDART	II a klasė
C022020000100	RADIX-ANKER-STANDART	II a klasė

Prod.nr.	Produkto žymėjimas	Produkto klasė
	RADIX-ANKER-STANDART	II a klasė
C022020000300	RADIX-ANKER-STANDART	II a klasė
C022020003B00	RADIX-ANKER-STANDART	II a klasė
C022500000000	STP ATKURIAMIEJI KAIŠČIAI (PINS)	II a klasė
C024100000800	Mooser Sistema	II a klasė
C024100001000	Mooser Sistema	II a klasė
C024300000000	Mooser Sistemos Integral rinkinys	II a klasė
C026100000100	Radix-Anker-Ilgas	II a klasė
C026100000200	Radix-Anker-Ilgas	II a klasė
C026100000300	Radix-Anker-Ilgas	II a klasė
C026320000100	Radix-Anker-Ilgas	II a klasė
C026320000200	Radix-Anker-Ilgas	II a klasė
C026320000300	Radix-Anker-Ilgas	II a klasė
C031000000000	Radix-Anker-Ilgas rinkinys	II a klasė
C032000000200	Radix-Anker-Standart Demo rinkinys	II a klasė
C032100020800	Unimetric T demo rinkinys	II a klasė
C032100021000	Unimetric T demo rinkinys	II a klasė
C032200000200	Radix-Anker-Ilgas Demo rinkinys	II a klasė
C032420000000	Radix-Anker-Ilgas	II a klasė
C055B00000100	MOOSER SISTEMA	II a klasė
C055B00000200	MOOSER SISTEMA	II a klasė
C055B00000300	MOOSER SISTEMA	II a klasė
C055B00000400	MOOSER SISTEMA	II a klasė
C055C00950800	MOOSER SISTEMA	II a klasė
C055C00951000	MOOSER SISTEMA	II a klasė
C055C01150800	MOOSER SISTEMA	II a klasė
C055C01151000	MOOSER SISTEMA	II a klasė
C055C01350800	MOOSER SISTEMA	II a klasė
C055C01351000	MOOSER SISTEMA	II a klasė
C055D00000800	MOOSER SISTEMA	II a klasė
C055D00001000	MOOSER SISTEMA	II a klasė
C055D01200800	MOOSER SISTEMA	II a klasė
C055D01400800	MOOSER SISTEMA	II a klasė
C055D01401000	MOOSER SISTEMA	II a klasė
C055D01601000	MOOSER SISTEMA	II a klasė
C060000000100	EasyPost	II a klasė
C060000000200	EasyPost	II a klasė
C060000000300	EasyPost	II a klasė
C060000000400	EasyPost	II a klasė
C060000000500	EasyPost	II a klasė
C060000000600	EasyPost	II a klasė
C060100000100	EasyPost	II a klasė
C060100000200	EasyPost	II a klasė
C060100000300	EasyPost	II a klasė
C060100000400	EasyPost	II a klasė
C060100000500	EasyPost	II a klasė
C060100000600	EasyPost	II a klasė
C060200800100	EasyPost	II a klasė
C060201000100	EasyPost	II a klasė
C060201300100	EasyPost	II a klasė
C061300000100	Radix Fiber Post	II a klasė
C061300000200	Radix Fiber Post	II a klasė
C061300000300	Radix Fiber Post	II a klasė
C061300000400	Radix Fiber Post	II a klasė
C061300000500	Radix Fiber Post	II a klasė
C061300000600	Radix Fiber Post	II a klasė
C061400800100	Radix Fiber Post Basic rinkinys	II a klasė

<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
C061400800200	Radix Fiber Post Trial rinkinys	II a klasė
C061401000100	Radix Fiber Post Basic rinkinys	II a klasė
C061401000200	Radix Fiber Post Trial rinkinys	II a klasė
C061401300100	Radix Fiber Post Basic rinkinys	II a klasė
C106KL0050A00	CYTCO-K	II a klasė
C106KL0050B00	CYTCO-K	II a klasė
C106KL0080A00	CYTCO-K	II a klasė
C106KL0080B00	CYTCO-K	II a klasė
C106KS0050A00	CYTCO-K	II a klasė
C107KL0050000	CYTCO-K	II a klasė
C107KL0080000	CYTCO-K	II a klasė
C107KS0050000	CYTCO-K	II a klasė
C108KL0050000	CYTCO-K	II a klasė
C108KL0080000	CYTCO-K	II a klasė
C108KS0050000	CYTCO-K	II a klasė
C117K00000000	CYTCO-K RINKINYS	II a klasė
C117KL0000100	CYTCO-K RINKINYS	II a klasė
C1187KL500100	CYTCO-K INTEGRAL RINKINYS	II a klasė
C1187KL500200	CYTCO-K INTEGRAL RINKINYS	II a klasė
C1187KL800100	CYTCO-K INTEGRAL RINKINYS	II a klasė
C1187KL800200	CYTCO-K INTEGRAL RINKINYS	II a klasė
C1187KS500100	CYTCO-K INTEGRAL RINKINYS	II a klasė
C193T00000100	RADIX-ANKER-STANDART	II a klasė
C193T00000200	RADIX-ANKER-STANDART	II a klasė
C193T00000300	RADIX-ANKER-STANDART	II a klasė
C193T00003B00	RADIX-ANKER-STANDART	II a klasė
C195C00000100	RADIX-ANKER-STANDART INTEGRAL RINKINYS	II a klasė
C195C00000200	RADIX-ANKER-STANDART INTEGRAL RINKINYS	II a klasė
C195C00000300	RADIX-ANKER-STANDART INTEGRAL RINKINYS	II a klasė
C195CT0000100	RADIX-ANKER-STANDART INTEGRAL RINKINYS	II a klasė
C195CT0000200	RADIX-ANKER-STANDART INTEGRAL RINKINYS	II a klasė
C195CT0000300	RADIX-ANKER-STANDART INTEGRAL RINKINYS	II a klasė
C195CT0003B00	RADIX-ANKER-STANDART INTEGRAL RINKINYS	II a klasė
C195CTR000200	RADIX-ANKER-STANDART INTEGRAL RINKINYS	II a klasė
C195CTR000300	RADIX-ANKER-STANDART INTEGRAL RINKINYS	II a klasė
C195CTR003B00	RADIX-ANKER-STANDART INTEGRAL RINKINYS	II a klasė
C195D00000000	RADIX-ANKER-STANDART RINKINYS	II a klasė
C195DT00000000	RADIX-ANKER-STANDART RINKINYS	II a klasė
C195ET00000000	RADIX-ANKER-STANDART RINKINYS	II a klasė
C211R00000000	STP ATKURIAMŲJŲ KAIŠČIŲ (PINS) INTEGRAL RINKINYS	II a klasė
C215TL0010800	UNIMETRIC X.X MM	II a klasė
C215TL0011000	UNIMETRIC X.X MM	II a klasė
C215TL0020800	UNIMETRIC X.X MM	II a klasė
C215TL0010800	UNIMETRIC X.X MM	II a klasė
C215TL0030000	UNIMETRIC X.X MM	II a klasė
C215TL0031000	UNIMETRIC X.X MM	II a klasė
C215TLS020800	UNIMETRIC X.X MM	II a klasė
C215TS0000800	UNIMETRIC X.X MM	II a klasė
C215TS0010800	UNIMETRIC X.X MM	II a klasė
C215TS0011000	UNIMETRIC X.X MM	II a klasė
C215TS0020800	UNIMETRIC X.X MM	II a klasė
C215TS0021000	UNIMETRIC X.X MM	II a klasė
C215TS0030800	UNIMETRIC X.X MM	II a klasė
C215TS0031000	UNIMETRIC X.X MM	II a klasė
C215U00010800	UNICLIP	II a klasė
C215U00011000	UNICLIP	II a klasė
C215U00020800	UNICLIP	II a klasė











<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
D006424002300	Plieno gražtai užbaigimo liepsnelės formos	II a klasė
D006424002500	Plieno gražtai užbaigimo liepsnelės formos	II a klasė
D006424002700	Plieno gražtai užbaigimo liepsnelės formos	II a klasė
D006424002900	Plieno gražtai užbaigimo liepsnelės formos	II a klasė
D006424003300	Plieno gražtai užbaigimo liepsnelės formos	II a klasė
D006424003500	Plieno gražtai užbaigimo liepsnelės formos	II a klasė
D006424003700	Plieno gražtai užbaigimo liepsnelės formos	II a klasė
D006424004000	Plieno gražtai užbaigimo liepsnelės formos	II a klasė
D020524000600	LN	II a klasė
D065B1401000	Plieno gražtai užbaigimo pumpuro formos	II a klasė
D065B1401800	Plieno gražtai užbaigimo pumpuro formos	II a klasė
D065B1402100	Plieno gražtai užbaigimo pumpuro formos	II a klasė
D065B1402700	Plieno gražtai užbaigimo pumpuro formos	II a klasė
D065B2401000	Plieno gražtai užbaigimo pumpuro formos	II a klasė
D065B2401200	Plieno gražtai užbaigimo pumpuro formos	II a klasė
D065B2401400	Plieno gražtai užbaigimo pumpuro formos	II a klasė
D065B2401600	Plieno gražtai užbaigimo pumpuro formos	II a klasė
D065B2401800	Plieno gražtai užbaigimo pumpuro formos	II a klasė
D065B2402100	Plieno gražtai užbaigimo pumpuro formos	II a klasė
D065B2402300	Plieno gražtai užbaigimo pumpuro formos	II a klasė
D065B2402900	Plieno gražtai užbaigimo pumpuro formos	II a klasė
D065B2404000	Plieno gražtai užbaigimo pumpuro formos	II a klasė
D171I14001200	Chirurginiai gražtai apvalūs, grubūs	II a klasė
D171I14001400	Chirurginiai gražtai apvalūs, grubūs	II a klasė
D171I14001600	Chirurginiai gražtai apvalūs, grubūs	II a klasė
D171I14001800	Chirurginiai gražtai apvalūs, grubūs	II a klasė
D171I14002100	Chirurginiai gražtai apvalūs, grubūs	II a klasė
D171I14002300	Chirurginiai gražtai apvalūs, grubūs	II a klasė
D171I14002500	Chirurginiai gražtai apvalūs, grubūs	II a klasė
D171I14002700	Chirurginiai gražtai apvalūs, grubūs	II a klasė
D171I14003100	Chirurginiai gražtai apvalūs, grubūs	II a klasė
D171I14003300	Chirurginiai gražtai apvalūs, grubūs	II a klasė
D171I14003500	Chirurginiai gražtai apvalūs, grubūs	II a klasė
D171I14003700	Chirurginiai gražtai apvalūs, grubūs	II a klasė
D171I14004000	Chirurginiai gražtai apvalūs, grubūs	II a klasė
D171I14004500	Chirurginiai gražtai apvalūs, grubūs	II a klasė
D171I14005000	Chirurginiai gražtai apvalūs, grubūs	II a klasė
D171I14005500	Chirurginiai gražtai apvalūs, grubūs	II a klasė
D171I14006000	Chirurginiai gražtai apvalūs, grubūs	II a klasė
D171I14006500	Chirurginiai gražtai apvalūs, grubūs	II a klasė
D172I14002100	Chirurginiai gražtai konuso formos, grubūs	II a klasė
D172I14002300	Chirurginiai gražtai konuso formos, grubūs	II a klasė
E012314000400	TC ertmės gražtai apvalūs	II a klasė
E012314000500	TC ertmės gražtai apvalūs	II a klasė
E012314000600	TC ertmės gražtai apvalūs	II a klasė
E012314000700	TC ertmės gražtai apvalūs	II a klasė
E012314000800	TC ertmės gražtai apvalūs	II a klasė
E012314000900	TC ertmės gražtai apvalūs	II a klasė
E012314001000	TC ertmės gražtai apvalūs	II a klasė
E012314001200	TC ertmės gražtai apvalūs	II a klasė
E012314001400	TC ertmės gražtai apvalūs	II a klasė
E012314001600	TC ertmės gražtai apvalūs	II a klasė
E012314001800	TC ertmės gražtai apvalūs	II a klasė
E012314002100	TC ertmės gražtai apvalūs	II a klasė
E012314002300	TC ertmės gražtai apvalūs	II a klasė
E012314002500	TC ertmės gražtai apvalūs	II a klasė
E012314002700	TC ertmės gražtai apvalūs	II a klasė







<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
E013837101200	TC ertmės gražtai konuso formos kampuoti	II a klasė
E013837101600	TC ertmės gražtai konuso formos kampuoti	II a klasė
E013837102100	TC ertmės gražtai konuso formos kampuoti	II a klasė
E014714000900	TC ertmės gražtai konuso formos apvalūs	II a klasė
E014734100900	TC ertmės gražtai konuso formos apvalūs	II a klasė
E014834101000	TC ertmės gražtai konuso formos apvalūs	II a klasė
E014834101200	TC ertmės gražtai konuso formos apvalūs	II a klasė
E014914001000	TC ertmės gražtai konuso formos apvalūs	II a klasė
E014914001200	TC ertmės gražtai konuso formos apvalūs	II a klasė
E014924101000	TC ertmės gražtai konuso formos apvalūs	II a klasė
E014924101400	TC ertmės gražtai konuso formos apvalūs	II a klasė
E014924101600	TC ertmės gražtai konuso formos apvalūs	II a klasė
E014934101000	TC ertmės gražtai konuso formos apvalūs	II a klasė
E014934101200	TC ertmės gražtai konuso formos apvalūs	II a klasė
E014934101400	TC ertmės gražtai konuso formos apvalūs	II a klasė
E014934101600	TC ertmės gražtai konuso formos apvalūs	II a klasė
E015114100000	Zekrya	II a klasė
E015115100000	Zekrya	II a klasė
E015124100000	Zekrya	II a klasė
E015134100000	Zekrya	II a klasė
E015135100000	Zekrya	II a klasė
E015224100000	Endo-Z	II a klasė
E015224100000	Endo-Z	II a klasė
E015237100000	Endo-Z	II a klasė
E015334101200	Transmetal	II a klasė
E015337101200	Transmetal	II a klasė
E015434100900	Transmetal	II a klasė
E015434101000	Transmetal	II a klasė
E015434101200	Transmetal	II a klasė
E015437101000	Transmetal	II a klasė
E015534101000	DC	II a klasė
E015534101200	DC	II a klasė
E015634101000	DC	II a klasė
E015634101200	DC	II a klasė
E015734100000	Day-Z	II a klasė
E015834101000	DC	II a klasė
E015834101200	DC	II a klasė
E020524100610	LN	II a klasė
E020524100800	LN	II a klasė
E020524100810	LN	II a klasė
E020524101010	LN	II a klasė
E020524101210	LN	II a klasė
E020524101410	LN	II a klasė
E020524101610	LN	II a klasė
E020524190000	LN	II a klasė
E051034100800	CF	II a klasė
E051034101000	CF	II a klasė
E051034101400	CF	II a klasė
E051134101400	CF	II a klasė
E051134101600	CF	II a klasė
E051234101800	CF	II a klasė
E051234102300	CF	II a klasė
E051534101200	CF	II a klasė
E051534101400	CF	II a klasė
E051734100900	CF	II a klasė
E051734101000	CF	II a klasė
E051734101200	CF	II a klasė

<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
E051834100700	CF	II a klasė
E051834100900	CF	II a klasė
E051834101000	CF	II a klasė
E051934101200	CF	II a klasė
E051934101400	CF	II a klasė
E051934101800	CF	II a klasė
E052034101000	CF	II a klasė
E052034101200	CF	II a klasė
E052134101000	CF	II a klasė
E052234101200	CF	II a klasė
E052234100900	CF	II a klasė
E052334101200	CF	II a klasė
E052434100900	CF	II a klasė
E052534101200	CF	II a klasė
E052634101000	CF	II a klasė
E052634101200	CF	II a klasė
E052734101400	CF	II a klasė
E052734101600	CF	II a klasė
E052834101000	CF	II a klasė
E052834101200	CF	II a klasė
E052834101400	CF	II a klasė
E052934101600	CF	II a klasė
E053034101400	CF	II a klasė
E053134101800	CF	II a klasė
E053234101200	CF	II a klasė
E053334101200	CF	II a klasė
E053334101400	CF	II a klasė
E053434101200	CF	II a klasė
E053434101400	CF	II a klasė
E053434101600	CF	II a klasė
E053434101600	CF	II a klasė
E053534101200	CF	II a klasė
E053634101400	CF	II a klasė
E053734101400	CF	II a klasė
E054014101600	CF	II a klasė
E054024101600	CF	II a klasė
E054035101600	CF	II a klasė
E054114101600	CF	II a klasė
E054124101600	CF	II a klasė
E054135101600	CF	II a klasė
E055034101100	CF	II a klasė
E055034102A00	CF	II a klasė
E055034102B00	CF	II a klasė
E055234101600	CF	II a klasė
E056035100000	Aryne-Z	II a klasė
E057034101200	DC	II a klasė
E057134101000	DC	II a klasė
E057134101200	DC	II a klasė
E058034101000	Transmetal	II a klasė
E059004100000	Metal-Master	II a klasė
E123A24101000	Excavabur	II a klasė
E123A24101200	Excavabur	II a klasė
E123A24101400	Excavabur	II a klasė
E123A24101600	Excavabur	II a klasė
E123A24101800	Excavabur	II a klasė
E123A24102100	Excavabur	II a klasė
E123A24102300	Excavabur	II a klasė

<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
E123A24102500	Excavabur	II a klasė
E123A24102700	Excavabur	II a klasė
E123A34101000	Excavabur	II a klasė
E123A34101200	Excavabur	II a klasė
E123A34101400	Excavabur	II a klasė
E123A34101600	Excavabur	II a klasė
E123A34101800	Excavabur	II a klasė
E123A34102100	Excavabur	II a klasė
E123A34102300	Excavabur	II a klasė
E123A34102500	Excavabur	II a klasė
E123A34102700	Excavabur	II a klasė
E129X14101000	TC ertmės grąžtai cilindro formos kampuotas	II a klasė
E129X14101200	TC ertmės grąžtai cilindro formos kampuotas	II a klasė
E129X34101000	TC ertmės grąžtai cilindro formos kampuotas	II a klasė
E129X34101200	TC ertmės grąžtai cilindro formos kampuotas	II a klasė
E129X34101400	TC ertmės grąžtai cilindro formos kampuotas	II a klasė
E130L34100900	TC ertmės grąžtai konuso formos kampuotas	II a klasė
E130L34101000	TC ertmės grąžtai konuso formos kampuotas	II a klasė
E130X14101200	TC ertmės grąžtai konuso formos kampuotas	II a klasė
E130X34101000	TC ertmės grąžtai konuso formos kampuotas	II a klasė
E130X34101200	TC ertmės grąžtai konuso formos kampuotas	II a klasė
E137X14101000	TC ertmės grąžtai cilindro formos kampuotas	II a klasė
E137X34101000	TC ertmės grąžtai cilindro formos kampuotas	II a klasė
E137X34101200	TC ertmės grąžtai cilindro formos kampuotas	II a klasė
E137X34101400	TC ertmės grąžtai cilindro formos kampuotas	II a klasė
E138L34100900	TC ertmės grąžtai konuso formos kampuotas	II a klasė
E138L34101000	TC ertmės grąžtai konuso formos kampuotas	II a klasė
E138L34101600	TC ertmės grąžtai konuso formos kampuotas	II a klasė
E138X34101000	TC ertmės grąžtai konuso formos kampuotas	II a klasė
E138X34101200	TC ertmės grąžtai konuso formos kampuotas	II a klasė
E150A14000600	TC amalgamos grąžtai	II a klasė
E150A14000800	TC amalgamos grąžtai	II a klasė
E150A24100800	TC amalgamos grąžtai	II a klasė
E150A24101000	TC amalgamos grąžtai	II a klasė
E150A24101200	TC amalgamos grąžtai	II a klasė
E150A32100800	TC amalgamos grąžtai	II a klasė
E150A32101000	TC amalgamos grąžtai	II a klasė
E150A34100600	TC amalgamos grąžtai	II a klasė
E150A34100800	TC amalgamos grąžtai	II a klasė
E150A34100900	TC amalgamos grąžtai	II a klasė
E150A34101000	TC amalgamos grąžtai	II a klasė
E150A34101200	TC amalgamos grąžtai	II a klasė
E150B24101000	TC amalgamos grąžtai	II a klasė
E150B24101200	TC amalgamos grąžtai	II a klasė
E150B24101400	TC amalgamos grąžtai	II a klasė
E150B34100800	TC amalgamos grąžtai	II a klasė
E150B34101000	TC amalgamos grąžtai	II a klasė
E150B34101200	TC amalgamos grąžtai	II a klasė
E150B34101400	TC amalgamos grąžtai	II a klasė
E150C34100800	TC amalgamos grąžtai	II a klasė
E150C34101400	TC amalgamos grąžtai	II a klasė
E510L34101400	CF	II a klasė
E536L34101400	CF	II a klasė
E537L34101400	CF	II a klasė
E541A14101600	CF	II a klasė
E541A24101600	CF	II a klasė
E541A35101600	CF	II a klasė

<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
F000133201000	Deimantiniai gražtai	II a klasė
F000133201400	Deimantiniai gražtai	II a klasė
F000134102300	Deimantiniai gražtai	II a klasė
F000134200900	Deimantiniai gražtai	II a klasė
F000134201000	Deimantiniai gražtai	II a klasė
F000134201200	Deimantiniai gražtai	II a klasė
F000134201400	Deimantiniai gražtai	II a klasė
F000134201600	Deimantiniai gražtai	II a klasė
F000134201800	Deimantiniai gražtai	II a klasė
F000134202300	Deimantiniai gražtai	II a klasė
F000134202500	Deimantiniai gražtai	II a klasė
F000134203300	Deimantiniai gražtai	II a klasė
F000134301000	Deimantiniai gražtai	II a klasė
F000134301400	Deimantiniai gražtai	II a klasė
F000134301600	Deimantiniai gražtai	II a klasė
F000134301800	Deimantiniai gražtai	II a klasė
F000134302300	Deimantiniai gražtai	II a klasė
F000134401200	Deimantiniai gražtai	II a klasė
F000134401400	Deimantiniai gražtai	II a klasė
F000134401600	Deimantiniai gražtai	II a klasė
F000134401800	Deimantiniai gražtai	II a klasė
F000134402300	Deimantiniai gražtai	II a klasė
F000234200900	Deimantiniai gražtai	II a klasė
F000234201000	Deimantiniai gražtai	II a klasė
F000234201200	Deimantiniai gražtai	II a klasė
F000234301200	Deimantiniai gražtai	II a klasė
F000234301400	Deimantiniai gražtai	II a klasė
F001034200900	Deimantiniai gražtai	II a klasė
F001034201000	Deimantiniai gražtai	II a klasė
F001034201200	Deimantiniai gražtai	II a klasė
F001034201400	Deimantiniai gražtai	II a klasė
F001034201600	Deimantiniai gražtai	II a klasė
F001034201800	Deimantiniai gražtai	II a klasė
F001034301400	Deimantiniai gražtai	II a klasė
F001934201200	Deimantiniai gražtai	II a klasė
F001934301000	Deimantiniai gražtai	II a klasė
F001934301200	Deimantiniai gražtai	II a klasė
F001934301400	Deimantiniai gražtai	II a klasė
F003234201400	Deimantiniai gražtai	II a klasė
F003234201600	Deimantiniai gražtai	II a klasė
F003334202700	Deimantiniai gražtai	II a klasė
F003834203300	Deimantiniai gražtai	II a klasė
F003934203700	Deimantiniai gražtai	II a klasė
F004134204500	Deimantiniai gražtai	II a klasė
F004234204500	Deimantiniai gražtai	II a klasė
F006834204000	Deimantiniai gražtai	II a klasė
F006834304000	Deimantiniai gražtai	II a klasė
F010733201000	Deimantiniai gražtai	II a klasė
F010733201200	Deimantiniai gražtai	II a klasė
F010734200800	Deimantiniai gražtai	II a klasė
F010734200900	Deimantiniai gražtai	II a klasė
F010734201000	Deimantiniai gražtai	II a klasė
F010734201200	Deimantiniai gražtai	II a klasė
F010734201400	Deimantiniai gražtai	II a klasė
F010734201600	Deimantiniai gražtai	II a klasė
F010734300900	Deimantiniai gražtai	II a klasė
F010734301000	Deimantiniai gražtai	II a klasė



<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
F016834202300	Deimantiniai gražtai	II a klasė
F016834202500	Deimantiniai gražtai	II a klasė
F017134101200	Deimantiniai gražtai	II a klasė
F017134201200	Deimantiniai gražtai	II a klasė
F017134201400	Deimantiniai gražtai	II a klasė
F017134301200	Deimantiniai gražtai	II a klasė
F017234201200	Deimantiniai gražtai	II a klasė
F017234201400	Deimantiniai gražtai	II a klasė
F017234201600	Deimantiniai gražtai	II a klasė
F017234301600	Deimantiniai gražtai	II a klasė
F017234301800	Deimantiniai gražtai	II a klasė
F017234401400	Deimantiniai gražtai	II a klasė
F017234401600	Deimantiniai gražtai	II a klasė
F017334101800	Deimantiniai gražtai	II a klasė
F017334201400	Deimantiniai gražtai	II a klasė
F017334201600	Deimantiniai gražtai	II a klasė
F017334201800	Deimantiniai gražtai	II a klasė
F017334301800	Deimantiniai gražtai	II a klasė
F017334401600	Deimantiniai gražtai	II a klasė
F019734200900	Deimantiniai gražtai	II a klasė
F019734202500	Deimantiniai gražtai	II a klasė
F019734301200	Deimantiniai gražtai	II a klasė
F019734301600	Deimantiniai gražtai	II a klasė
F019734301800	Deimantiniai gražtai	II a klasė
F019734402500	Deimantiniai gražtai	II a klasė
F019834101600	Deimantiniai gražtai	II a klasė
F019834101800	Deimantiniai gražtai	II a klasė
F019834102300	Deimantiniai gražtai	II a klasė
F019834201600	Deimantiniai gražtai	II a klasė
F019834301200	Deimantiniai gražtai	II a klasė
F019834301400	Deimantiniai gražtai	II a klasė
F019834301600	Deimantiniai gražtai	II a klasė
F019834301800	Deimantiniai gražtai	II a klasė
F019834302300	Deimantiniai gražtai	II a klasė
F019834401400	Deimantiniai gražtai	II a klasė
F019834401600	Deimantiniai gražtai	II a klasė
F019834401800	Deimantiniai gražtai	II a klasė
F019934201200	Deimantiniai gražtai	II a klasė
F019934201400	Deimantiniai gražtai	II a klasė
F019934201600	Deimantiniai gražtai	II a klasė
F019934201800	Deimantiniai gražtai	II a klasė
F019934202300	Deimantiniai gražtai	II a klasė
F019934301400	Deimantiniai gražtai	II a klasė
F019934401600	Deimantiniai gražtai	II a klasė
F019934401800	Deimantiniai gražtai	II a klasė
F019937401600	Deimantiniai gražtai	II a klasė
F021934101200	Deimantiniai gražtai	II a klasė
F021934201200	Deimantiniai gražtai	II a klasė
F021934201600	Deimantiniai gražtai	II a klasė
F022034201400	Deimantiniai gražtai	II a klasė
F022534201200	Deimantiniai gražtai	II a klasė
F022534201600	Deimantiniai gražtai	II a klasė
F023334200900	Deimantiniai gražtai	II a klasė
F023334201000	Deimantiniai gražtai	II a klasė
F023334201200	Deimantiniai gražtai	II a klasė
F023334201400	Deimantiniai gražtai	II a klasė
F023334401200	Deimantiniai gražtai	II a klasė

<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
F023434201200	Deimantiniai gražtai	II a klasė
F023434301200	Deimantiniai gražtai	II a klasė
F023434301400	Deimantiniai gražtai	II a klasė
F023434301600	Deimantiniai gražtai	II a klasė
F023434301800	Deimantiniai gražtai	II a klasė
F024534301000	Deimantiniai gražtai	II a klasė
F024534301400	Deimantiniai gražtai	II a klasė
F024934201200	Deimantiniai gražtai	II a klasė
F024934001200	Deimantiniai gražtai	II a klasė
F024934001400	Deimantiniai gražtai	II a klasė
F024934001600	Deimantiniai gražtai	II a klasė
F024934101200	Deimantiniai gražtai	II a klasė
F024934101400	Deimantiniai gražtai	II a klasė
F024934201000	Deimantiniai gražtai	II a klasė
F024934301200	Deimantiniai gražtai	II a klasė
F024934301400	Deimantiniai gražtai	II a klasė
F024934301600	Deimantiniai gražtai	II a klasė
F024934401200	Deimantiniai gražtai	II a klasė
F024934401400	Deimantiniai gražtai	II a klasė
F025034001200	Deimantiniai gražtai	II a klasė
F025034101200	Deimantiniai gražtai	II a klasė
F025034101600	Deimantiniai gražtai	II a klasė
F025034201200	Deimantiniai gražtai	II a klasė
F025034301200	Deimantiniai gražtai	II a klasė
F025034301600	Deimantiniai gražtai	II a klasė
F025734001600	Deimantiniai gražtai	II a klasė
F025734002300	Deimantiniai gražtai	II a klasė
F025734101600	Deimantiniai gražtai	II a klasė
F025734102300	Deimantiniai gražtai	II a klasė
F025734301600	Deimantiniai gražtai	II a klasė
F025734302300	Deimantiniai gražtai	II a klasė
F027734002300	Deimantiniai gražtai	II a klasė
F027734101800	Deimantiniai gražtai	II a klasė
F027734201800	Deimantiniai gražtai	II a klasė
F027734202300	Deimantiniai gražtai	II a klasė
F027734302300	Deimantiniai gražtai	II a klasė
F028834001000	Deimantiniai gražtai	II a klasė
F028834200900	Deimantiniai gražtai	II a klasė
F028834201000	Deimantiniai gražtai	II a klasė
F028834201200	Deimantiniai gražtai	II a klasė
F028834201400	Deimantiniai gražtai	II a klasė
F028834301000	Deimantiniai gražtai	II a klasė
F028834301200	Deimantiniai gražtai	II a klasė
F028934001000	Deimantiniai gražtai	II a klasė
F028934101400	Deimantiniai gražtai	II a klasė
F028934201000	Deimantiniai gražtai	II a klasė
F028934201200	Deimantiniai gražtai	II a klasė
F028934201400	Deimantiniai gražtai	II a klasė
F028934301200	Deimantiniai gražtai	II a klasė
F028934301400	Deimantiniai gražtai	II a klasė
F028934301600	Deimantiniai gražtai	II a klasė
F028934401400	Deimantiniai gražtai	II a klasė
F029034001000	Deimantiniai gražtai	II a klasė
F029034101200	Deimantiniai gražtai	II a klasė
F029034101400	Deimantiniai gražtai	II a klasė
F029034201400	Deimantiniai gražtai	II a klasė
F029034301200	Deimantiniai gražtai	II a klasė

<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
F029034301400	Deimantiniai gražtai	II a klasė
F029034301600	Deimantiniai gražtai	II a klasė
F029734301200	Deimantiniai gražtai	II a klasė
F029734301400	Deimantiniai gražtai	II a klasė
F029734301600	Deimantiniai gražtai	II a klasė
F029834101600	Deimantiniai gražtai	II a klasė
F029834201200	Deimantiniai gražtai	II a klasė
F029834201600	Deimantiniai gražtai	II a klasė
F029834301200	Deimantiniai gražtai	II a klasė
F029834301400	Deimantiniai gražtai	II a klasė
F029834301600	Deimantiniai gražtai	II a klasė
F029834301800	Deimantiniai gražtai	II a klasė
F029834302100	Deimantiniai gražtai	II a klasė
F029834401600	Deimantiniai gražtai	II a klasė
F029934101600	Deimantiniai gražtai	II a klasė
F029934101800	Deimantiniai gražtai	II a klasė
F029934201600	Deimantiniai gražtai	II a klasė
F029934301200	Deimantiniai gražtai	II a klasė
F029934301400	Deimantiniai gražtai	II a klasė
F029934301600	Deimantiniai gražtai	II a klasė
F029934301800	Deimantiniai gražtai	II a klasė
F029934302100	Deimantiniai gražtai	II a klasė
F029934401600	Deimantiniai gražtai	II a klasė
F030434201600	Deimantiniai gražtai	II a klasė
F030434202300	Deimantiniai gražtai	II a klasė
F049634201200	Deimantiniai gražtai	II a klasė
F054033200900	Deimantiniai gražtai	II a klasė
F054034100900	Deimantiniai gražtai	II a klasė
F054034101000	Deimantiniai gražtai	II a klasė
F054034200900	Deimantiniai gražtai	II a klasė
F054034300900	Deimantiniai gražtai	II a klasė
F054034301000	Deimantiniai gražtai	II a klasė
F054434201600	Deimantiniai gražtai	II a klasė
F054534201600	Deimantiniai gražtai	II a klasė
F054634301600	Deimantiniai gražtai	II a klasė
F054634301800	Deimantiniai gražtai	II a klasė
F069735301600	Deimantiniai gražtai	II a klasė
F069735301800	Deimantiniai gražtai	II a klasė
F095100000300	Izometriniai gražtai	II a klasė
F095100000400	Izometriniai gražtai	II a klasė
F095134000300	Izometriniai gražtai	II a klasė
F095134000400	Izometriniai gražtai	II a klasė
F095134090000	Izometriniai gražtai	II a klasė
F095134100100	Izometriniai gražtai	II a klasė
F095134100200	Izometriniai gražtai	II a klasė
F095134100300	Izometriniai gražtai	II a klasė
F095134100400	Izometriniai gražtai	II a klasė
F095134190000	Izometriniai gražtai	II a klasė
F095134200100	Izometriniai gražtai	II a klasė
F095134200200	Izometriniai gražtai	II a klasė
F095134200300	Izometriniai gražtai	II a klasė
F095134200400	Izometriniai gražtai	II a klasė
F095230090000	Izometriniai gražtai	II a klasė
F095234000100	Izometriniai gražtai	II a klasė
F095234000200	Izometriniai gražtai	II a klasė
F095333001400	Deimantiniai gražtai	II a klasė
F0953334001400	Deimantiniai gražtai	II a klasė

<b>Prod.nr.</b>	<b>Produkto žymėjimas</b>	<b>Produkto klasė</b>
F095335001400	Deimantiniai gražtai	II a klasė
F116100000000	Gingikit rinkinys	II a klasė
F116134210100	Gingikit rinkinys	II a klasė
F116134210200	Gingikit rinkinys	II a klasė
F116134210300	Gingikit rinkinys	II a klasė
F116135210400	Gingikit rinkinys	II a klasė
F116135210500	Gingikit rinkinys	II a klasė
F116135210600	Gingikit rinkinys	II a klasė
F116136210700	Gingikit rinkinys	II a klasė
F116136210800	Gingikit rinkinys	II a klasė
F116136210900	Gingikit rinkinys	II a klasė
F116200000000	Ceramprep rinkinys	II a klasė
F130000000000	Bogopolsky rinkinys	II a klasė
S209000310100	X-Post	II a klasė
S209000310200	X-Post	II a klasė
S209000310300	X-Post	II a klasė
S209000310400	X-Post	II a klasė

215-08-05, Ballaigues

Luc Etienne  
Valdymo atstovas

# EC Certificate - Production Quality Assurance

Directive 93/42/EEC on Medical Devices, Annex V

**No.** CE 00847  
**Issued To:** **Kerr Corporation,  
Also Doing Business as  
Pentron Clinical  
1717 West Collins Avenue  
Orange  
California 92867  
USA**

In respect of:

**The manufacture of dental composite restorative materials and accessories (ribbons, core materials, etchants, modifiers, primers), dental cements, dental sealants, fibre posts and associated non-sterile dental instruments for attachment to an active device.**

on the basis of our examination of the quality assurance system under the requirements of Council Directive 93/42/EEC, Annex V. The quality assurance system meets the requirements of the directive. For the placing on the market of class IIb and class III products an Annex III certificate is required.

For and on behalf of BSI, a Notified Body for the above Directive (Notified Body Number 0086):



Gary Fenton, Global Assurance Director

First Issued: **29 August 1995**

Date: **27 August 2014**

Expiry Date: **28 August 2019**

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Page 1 of 1

Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the required surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI.

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

# EC Certificate - Production Quality Assurance

Directive 93/42/EEC on Medical Devices, Annex V

## List of Significant Subcontractors

Recognised as being involved in services relating to the product covered by:

Certificate No: **CE 00847**  
Date: **27 August 2014**  
Issued To: **Kerr Corporation,  
Also Doing Business as  
Pentron Clinical  
1717 West Collins Avenue  
Orange  
California 92867  
USA**

<b>Subcontractor:</b>	<b>Service(s) supplied</b>
Kerr Italia S.r.l. Via Passanti, 332 Scafati (SA) 84018 Italy	<b>EU Representative</b>
SDS de Mexico S. de R.L. de C.V., A Subsidiary of Ormco Corporation Circuito Sur No. 31 Parque Industrial Nelson Mexicali Baja California C.P.21395 Mexico	<b>Packaging</b>
SpofaDental a.s Markova 238 Jicín CZ-506 01 Czech Republic	<b>EU Representative</b>

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# EC Certificate - Production Quality Assurance Certificate History

Certificate No: **CE 00847**  
 Date: **27 August 2014**  
 Issued To: **Kerr Corporation,  
 Also Doing Business as  
 Pentron Clinical  
 1717 West Collins Avenue  
 Orange  
 California 92867  
 USA**

Date	Reference Number	Action
29 August 1995	-	First issued
05 November 1997	-	Address change
19 November 2004	-	Revised wording of scope from 'dental composite restorative materials' to 'dental composite restorative systems'. Certificate renewal and reissue in new format.
05 July 2006	-	Correction to company name from Kerr Dental Materials Centre to Kerr Corporation
07 September 2009	7437306	Addition of 'Kerr Italia, SpA' as EU Representative Certificate renewal
05 November 2009	7452138	Company name changed, addition of "Also Doing Business as Pentron Clinical" Addition of 3 new sub contractors, "Pentron Clinical Technologies, LLC" for manufacturing, "SDS de Mexico SA de C.V" for packaging, and CEpartner4U for EU Rep. sub contractor activities Changed the subcontractor name for the Kerr EU rep from Kerr Italia Spa to Kerr Italia S.r.l.
26 October 2010	7596167	Clarification of scope and removal of the subcontractor Pentron Clinical Technologies, LLC, 68 North Plains Industrial Road, Wallingford, CT 06492, USA

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Page 1 of 2

Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the required surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI.  
 This certificate was issued electronically and is bound by the conditions of the contract.

# EC Certificate - Production Quality Assurance Certificate History

Certificate No: **CE 00847**  
 Date: **27 August 2014**  
 Issued To: **Kerr Corporation,  
 Also Doing Business as  
 Pentron Clinical  
 1717 West Collins Avenue  
 Orange  
 California 92867  
 USA**

Date	Reference Number	Action
27 August 2014	8166389	Certificate Renewal, removal of CEPartner4U and addition of "SpofaDental" as EU representative for Pentron product lines.

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Page 2 of 2

Validity of this certificate is conditional on the quality system being maintained to the requirements of the Directive as demonstrated through the required surveillance activities of the Notified Body. This approval excludes all products designed and/or manufactured by a third party on behalf of the company named on this certificate, unless specifically agreed with BSI.

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

**EB Sertifikatas – Gamybos kokybės užtikrinimas**

Medicinos prietaisų 93/42/EEB Direktyva, V priedas

Nr. **CE 00847**  
Išduota: **Kerr Corporation**  
**Taip pat vykdo verslą, kaip**  
**Petron Clinical**  
**1717 West Collins aveniu**  
**Orange**  
**California 92867**  
**USA (JAV)**

Dėl:

**Odontologinių kompozicinių restauracinių medžiagų ir priedų gamyba (juostelės, kulties medžiagos, ėsdintojai, modifikatoriai, praimeriai), odontologiniai cementai, odontologiniai silantai, stiklo pluošto kaiščiai ir susiję nesterilūs odontologiniai instrumentai priskiriami prie aktyvaus prietaiso.**

remiantis mūsų kokybės užtikrinimo sistemos tyrimu pagal Tarybos direktyvos 93/42/EEB V priedo reikalavimus. Kokybės užtikrinimo sistema atitinka direktyvos reikalavimus. Pateikiant rinkai IIb ir III klasės produktus reikalingas sertifikato III priedas.

BSI vardu, Notifikavimo įstaigos viršuje paminėtos direktyvos (Notifikavimo įstaigos numeris 0086):

Gary Fenton, Globalus užtikrinimo direktorius

Pirmas leidimas: **1995, rugpjūčio 29**Data: **2014 rugpjūčio 27**Galiojimo data: **2014 rugpjūčio 27**

1 psl.

VERTIMAS TEISINGAS

UAB „Plandent“ direktorius Andrius Preišegalavičius



**EB Sertifikatas – Gamybos kokybės užtikrinimas**

Medicinos prietaisų 93/42/EEB Direktyva, V priedas

**Svarbių subrangovų sąrašas**

Pripažinta kaip įtraukta į paslaugas, susijusias su produkto, kuriam taikoma:

Sertifikato Nr. **CE 00847**  
Data: **2014 rugpjūčio 27**  
Išduota: **Kerr Corporation**  
**Taip pat vykdo verslą, kaip**  
**Petron Clinical**  
**1717 West Collins aveniu**  
**Orange**  
**California 92867**  
**USA (JAV)**

**Subrangovas:****Teikiamos paslauga(-os)**

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Kerr Italia S.r.l.  
Via Passanti, 332  
Scafati (SA)  
84018  
Italy (Italija)

**ES atstovas**

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SDS de Mexico S. de R.L. de C.V.,  
ASubsidiary of Ormco Corporation  
Circuito Sur No. 31  
Parque Industrial Nelson  
Mexicali  
Baja California  
C.P.21395  
Mexico (Meksika)

**Pakuotė**

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SpofaDental a.s.  
Markova 238  
Jicin  
CZ-506 01  
Czech Republik (Čekijos respublika)

**ES atstovas**

## EB Sertifikatas – Gamybos kokybės užtikrinimas Sertifikato istorija

Sertifikato Nr. **CE 00847**  
 Data: **2014 rugpjūčio 27**  
 Išduota: **Kerr Corporation**  
**Taip pat vykdo verslą, kaip**  
**Petron Clinical**  
**1717 West Collins aveniu**  
**Orange**  
**California 92867**  
**USA (JAV)**

Data	Nuorodos numeris	Veikla
1995 rugpjūčio 29	-	Pirmas leidimas
1997 lapkričio 5	-	Pakeistas adresas
2004 Lapkričio 19	-	Peržiūrėta redakcija taikymo srities iš „dantų kompozicinės restauravimo medžiagos“ į „dantų kompozicinės restauravimo sistemas“. Sertifikato atnaujinimas ir perleidimas nauju formatu.
2006 liepos 5	-	Pataisa įmonės pavadinimo iš Kerr Dental Materials Centre į Kerr Corporation
2009 rugsėjo 7	7437306	Papildymas „Kerr Italia SpA“ ES atstovas Sertifikato atnaujinimas
2009 lapkričio 5	7452138	Kompanijos pavadinimo keitimas, papildomai „Taip pat vykdo verslą, kaip Petron Clinical“ Papildomai 3 nauji subrangovai, „Petron Clinical Technologies, LLC“ gamyba, „SDS de Mexico SA de C.V“ pakuotė ir CEPartner4U ES resp. subrangovų veikla Subrangovo pavadinimo keitimas Kerr EU rep iš Kerr Italia Spa į Kerr Italia S.r.l.
2010 spalio 26	7596167	Taikymo srities patikslinimas ir subrangovo Pentron Clinical Technologies, LLC, 68 North Plains Industrial Road, Wallingford, CT 06492, USA pašalinimas



## EB Sertifikatas – Gamybos kokybės užtikrinimas Sertifikato istorija

Sertifikato Nr. **CE 00847**  
Data: **2014 rugpjūčio 27**  
Išduota: **Kerr Corporation**  
**Taip pat vykdo verslą, kaip**  
**Petron Clinical**  
**1717 West Collins aveniu**  
**Orange**  
**California 92867**  
**USA (JAV)**

Data	Nuorodos numeris	Veikla
2014 rugpjūčio 27	8166389	Sertifikato atnaujinimas, CEPartner4U pašalinimas ir „SpofaDental“ kaip EB atstovo Pentron produktų linijų, papildymas.

2 psl. iš 2

  
VERTIMAS TEISINGAS  
  
UAB „Plandent“ direktorius Andrius Preišegalavičius

## EC – Declaration of Conformity

By virtue of this declaration of conformity,  
KerrHawe SA  
ensures and declares  
that the following items are Class I products  
and are manufactured in conformity with the provisions  
of the Council Directive 93/42/EEC and the  
technical documentation referred to in Annex VII, Section 3 and Annex V  
as amended by 2007/47/EC,  
thus, meeting the requirements of the Medical Device Directive

„X-Rays Film and Sensor Holder“ Class I,  
classified as „Guidance Devices “ intended for transient uses,  
containing the products indicated on the enclosed list „RB“

  
Alessandro Bressan  
(Managing Director Operations Europe)

20.04.2011  
Date

  
Greet Pfister  
(Quality Assurance Manager)

31.03.11  
Date

KerrHawe SA  
Via Strecce 4, 6934 Bioggio, Switzerland

**List RB**

The table below indicates the semi-finished goods and the relative finished product items.

Article Name	Semi Finished Article Number	Finished Products Article Number
Kwik-Bite	210030	270
Index EUR	210039	670
		671
		672
Index with ring	210045	1020
		1021
		1022
		1720
		1721
		1770
		1780
		1786
		1790
		1791
		1792
		2700
		2720
		2800
		2801
2802		
2900		
2901		
2902		
Parobite	210054	1720
		1770
SuperBite anterior "plus"	260008	1020
		1021
		1720
		670
		671
SuperBite posterior "plus"	260009	1020
		1022
		1720
		670
		672
Kwik Bite Senso assemblato	260010	2700
		2720
Kwik Bite Plus	260017	1720
		1780

Article Name	Semi Finished Article Number	Finished Products Article Number
Super Bite Senso Anterior	260018	2720 2800 2801
Super Bite Senso Posterior	260019	2720 2800 2802
Endo Bite Senso Anterior	260020	2720 2900 2901
Endo Bite Anterior Plus	26021	2720 2900 2902
Endo Bite Anterior Plus	260022	1720 1790 1791
Endo Bite Posterior Plus	260023	1720 1790 1792

## ES Atitikties deklaracija

Remiantis šia atitikties deklaracija  
KerrHawe SA  
garantuoja ir pareiškia, kad sekančios prekės yra I klasės produktai  
Ir yra pagaminti bei atitinka  
Tarybos direktyvos 93/42/EEC reikalavimus  
Pagal techninę dokumentaciją atitinkamai VII priedo, 3 dalies ir V priedo,  
„Tarpdantiniai kaiščiai“ I klasė,  
Klasifikuojamos kaip „Anatominiai modeliuotojai“ skirti laikinam (tarpiniam) naudojimui,  
Apimantys produktus nurodytus priede.

Gert Silber  
Generalinis direktorius Europos sk.

Greet Pfister  
Kokybės valdymo direktorius

Kerr Hawe sa  
P.O.Box. 268, 6934 Bioggio, Šveicarija

### Pastų sąrašas

Apačioje esančioje lentelėje nurodyti pusgaminiai ir tarpusavyje susiję produktai.

Pavadinimas	Artik.Nr.	Produkto Nr.
Mediniai kaištukai 05 turkio spalvos	200009	101, 102, 1600, 1620, 2008, 2080, 2090, 822, 824, 828, 829
Mediniai kaištukai 10 oranžinės spalvos	200012	101, 102, 103, 104, 1500, 1600, 1620, 2008, 2080, 2090, 630, 718, 725, 785, 786, 790, 822, 823, 824, 826, 827, 828, 829, 830
Mediniai kaištukai 10 žalios spalvos PL	200013	2422, 2422/10, 2423, 2427, 2428, 829
Mediniai kaištukai 20 baltos spalvos	200023	101, 102, 103, 104, 1500, 1600, 1620, 2008, 2080, 2090, 630, 785, 786, 822, 824, 826, 827, 828, 829, 830
Mediniai kaištukai 20 raudonos spalvos PL	200024	2422, 2422/20, 2423, 2427, 2428, 829
Mediniai kaištukai 20 baltos spalvos	200025	829
Mediniai kaištukai 30 žalios spalvos	200032	101, 102, 103, 104, 1500, 1600, 1620, 630, 718, 725, 785, 786, 790, 822, 823, 824, 826, 827, 828, 829, 830
Mediniai kaištukai 30 baltos spalvos PL	200033	2422, 2422/30, 2423, 2427, 2428, 829
Mediniai kaištukai 40 geltonos spalvos	200043	101, 102, 103, 104,

PL		1500, 822, 823, 824, 826, 827, 828, 829
Mediniai kaištukai 40 oranžinės spalvos PL	200044	2422, 2422/40, 2423, 2427, 2428, 829
Mediniai kaištukai 50 mėsvos spalvos	200053	1500, 1600, 1620, 630, 718, 725, 822, 823, 824, 826, 827, 828, 829, 830
Mediniai kaištukai 60 raudonos spalvos	200063	1500, 822, 823, 826, 827, 828, 829
Mediniai kaištukai 70 baltos spalvos	200072	1500, 822, 823, 827, 829
Mediniai kaištukai 40 geltonos spalvos	200073	829
Mediniai kaištukai 50 mėsvos spalvos	200074	829
Mediniai kaištukai 60 raudonos spalvos	200075	829
Luciwedges soft 1.0	211042	101, 104, 1500, 1600, 2070, 2071, 2072, 700, 710, 770, 780, 785, 786, 790
Luciwedges soft 1.2	211043	101, 1500, 1600, 2070, 2071, 2072, 700, 771, 780, 785, 790
Luciwedges soft 1.6	211044	101, 104, 1500, 1600, 2070, 2071, 2072, 700, 710, 772, 773, 780, 785, 786, 790
Adapt Luciwedges soft 1.3	211133	101, 102, 105, 1620, 730, 731, 750, 780, 785
Adapt Luciwedges soft 1.7	211134	101, 102, 1620, 730, 732, 750, 780, 785
Adapt Luciwedges soft 2.1	211135	101, 102, 1620, 730, 733, 750, 780, 785
Adapt Luciwedges soft 1.1	211173	734



  
 VERTIMAS TEISINGAS  
 UAB „Plandent“ direktorius Andrius Preišegalavičius

## EB Atitikties deklaracija

Remiantis šia atitikties deklaracija  
KerrHawe SA  
garantuoja ir pareiškia, kad sekančios prekės yra Ila klasės produktai  
Ir yra pagaminti bei atitinka  
Tarybos direktyvos 93/42/EEB reikalavimus  
pagal techninę dokumentaciją atitinkamai VII priedo, 3 dalies ir V priedo,  
kaip nurodyta 2007/47/EB,  
ir atitinka medicinos prietaisų direktyvos reikalavimus

„Rentgeno juostelių ir jutiklio laikiklis“ I klasė,  
klasifikuojami kaip „Nurodomieji prietaisai“ skirti laikinam (tarpiniam) naudojimui,  
apimantys produktus nurodytus priede „RB“.

Alessandro Bressanno      2011-04-20  
Europos sk. Veiklos vadovas      Data

Greet Pfister  
Kokybės valdymo direktorius

Kerr Hawe sa  
Via Strecce 4, 6934 Bioggio, Šveicarija

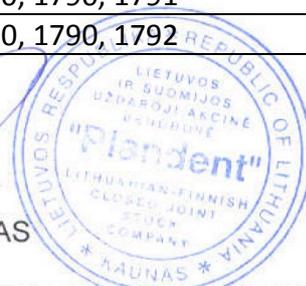
### RB sąrašas

Apačioje esančioje lentelėje nurodyti pusgaminiai ir tarpusavyje susiję produktai.

Pavadinimas	Artik.Nr.	Produkto Nr.
Kwik-Bite	210030	270
Index EUR	210039	670, 671, 672
Index su žiedu	210045	1020, 1021, 1022, 1720, 1721, 1770, 1780, 1786, 1790, 1791, 1792, 2700, 2720, 2800, 2801, 2802, 2900, 2901, 2901
Parobite	210054	1720, 1770
SuperBite anterior "plus"	260008	1020, 1021, 1720, 670, 671
SuperBite posterior "plus"	260009	1020, 1022, 1720, 670, 672
Kwik Bite Senso rinkinys	260010	2700, 2720,
Kwik Bite Plus	260017	1720, 1780
Super Bite Senso anterior	260018	2720, 2800, 2801
Super Bite Senso posterior	260019	2720, 2800, 2802
Endo Bite Senso anterior	260020	2720, 2900, 2901
Endo Bite anterior plus	26021	2720, 2900, 2902
Endo Bite anterior plus	260022	1720, 1790, 1791
Endo Bite posterior plus	260023	1720, 1790, 1792

VERTIMAS TEISINGAS

UAB „Plandent“ direktorius Andrius Preišgalavičius



## ES Atitikties deklaracija

Remiantis šia atitikties deklaracija  
KerrHawe SA  
garantuoja ir pareiškia, kad sekančios prekės yra I klasės produktai  
Ir yra pagaminti bei atitinka  
Tarybos direktyvos 93/42/EEC reikalavimus  
Pagal techninę dokumentaciją atitinkamai VII priedo, 3 dalies ir V priedo,

„Juostelės ir matricinės juostelės“ I klasė,  
Klasifikuojamos kaip „Anatominės formos“ skirti laikinam (tarpiniam) naudojimui,  
Apimantys produktus nurodytus priede „Juostelės“.

Annunziata Schettino  
Gamybos vadovas Europos sk.

Greet Pfister  
Kokybės valdymo direktorius

Kerr Hawe sa  
P.O.Box. 268, 6934 Bioggio, Šveicarija

### Pastų sąrašas

Apačioje esančioje lentelėje nurodyti pusgaminiai ir tarpusavyje susiję produktai.

Artik.Nr.	Tarpinis Art Nr.	Produkto Nr.
Hawe Striproll (juostelė) 6 mm plati 15Mt	114020	685
Hawe Striproll (juostelė) 8 mm plati 15Mt	114021	686
Hawe Striproll (juostelė) 10 mm plati 15Mt	114022	687
Nastro PP X skaidri juostelė 6 mm geltona	114044	695
Hawe skaidrios juostelės raudona 8 mm lenkta	114045	696
Hawe skaidrios juostelės mėlyna 10 mm lenkta	114046	697
Hawe skaidrios juostelės raudona 8 mm	114048	101, 102, 1620, 640, 641, 650, 651, 691
Hawe skaidrios juostelės mėlyna 10 mm	114049	101, 102, 1620, 640, 642, 650, 652, 692
Hawe skaidrios juostelės mėlyna 0.05x6 mm	114053	2685
Hawe skaidrios juostelės mėlyna 0.05x8 mm	114054	2686
Hawe skaidrios juostelės mėlyna 0.05x10 mm	114055	2687

  
VERTIMAS TEISINGAS  


UAB „Plandent“ direktorius Andrius Preišegalavičius



KaVo. Dental Excellence.

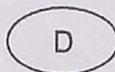
EG-Konformitätserklärung  
EC-Declaration of conformity  
CE-Déclaration de conformité  
Dichiarazione di conformità con le norme CE  
Prohlášení o shodě s normami ES  
Declaración de Conformidad de la CE

**Manufacturer:**  
**Kaltenbach & Voigt GmbH**  
**Bismarckring 39**  
**D-88400 Biberach**

**PiezoLED®**

**93 / 42 / EEC**  
**EN 1639 : 2009**

**Medical Devices / Annex I**  
**Dentistry - Instruments**



Wir erklären, dass die von uns hergestellten Produkte auf die sich diese Erklärung bezieht, mit den grundlegenden Anforderungen gemäß den Bestimmungen der obigen Richtlinie übereinstimmen. Zur Beurteilung des Ergebnisses wurden die benannten Normen oder normative Dokumente angewandt.



We declare that the products manufactured by us to which this statement refers, conform to the essential requirements according to the above-mentioned directive. The specified standards or normative documents were applied to evaluate the product.



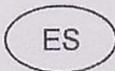
Nous déclarons que les produits que nous fabriquons, auxquels se réfère cette déclaration, sont conformes aux exigences essentielles selon les dispositions de la directive susmentionnée. Les normes ou autres documents normatifs désignés sont utilisés pour le jugement de ce produit.



Dichiariamo che i prodotti da noi fabbricati, a cui si riferisce la presente dichiarazione, sono conformi ai requisiti fondamentali ai sensi delle disposizioni della direttiva summenzionata. Per la valutazione del prodotto sono stati applicati le norme o i documenti normativi citati.



Prohlašujeme, že se námi vyráběné výrobky, na které se vztahuje toto prohlášení, shodují se základními požadavky podle předpisů výše uvedené směrnice. K posouzení výrobku byly použity uvedené normy nebo normativy.



Por la presente declaramos que los productos que nosotros fabricamos, y a los que hace referencia esta declaración, cumplen los requisitos básicos de conformidad con las disposiciones de la directiva arriba mencionada. Las normas o documentos normativos mencionados se aplican en la evaluación de dichos productos.



Biberach  
effective date 02.10.2010  
expiry date 01.10.2017



Certification GmbH  
Handwerkstrasse 15  
D-70565 Stuttgart

Norbert Glaeske  
Quality Director



Pardavimo vadybininkas  
Laimonas Ženkus

Tomas Kiškauskas  
Med. inžinierius

Vadybininkė  
Vaida Pečiulytė

/Logotipas: KaVo. Dental Excellence./

EG-Konformitätseklärung  
 EB atitikties deklaracija  
 CE-Déclaration de conformité  
 Dichiarazione di conformità con le norme CE  
 Prohlášení o shodě s normami ES  
 Declaracion de Conformidad de la CE

**Gamintojas:**

**Kaltenbach & Voigt GmbH**  
**Bismarckring 39,**  
**88400 Biberach**

**PiezoLED®**

**93 / 42 / EEB    Medicinos prietaisai / I priedas**  
**EN 1639 : 2009    Odontologijos instrumentai**

D                    Wir erklären, dass die von uns hergestellter Produkte auf die sich diese Erklärung bezieht. mit den grundlegenden Anforderungen gemäß den Bestimmungen der obigen Richtlinie übereinstimmen. Zur Beurteilung des Erzeugnisses wurden die benannten Normen oder normatyve Dokumente angewandt.

GB                  Patvirtiname, kad šioje deklaracijoje aprašyti mūsų gaminami gaminiai atitinka esminius aukščiau minėtos direktyvos reikalavimus. Produkto įvertinimui buvo taikomi atitinkami standartai ir norminiai dokumentai.

FR                  Nous déclarons que les produits que nous fabriquons, auxquels se réfère cette déclaration, sont conformes aux exigences essentielles selon les dispositions de la directive susmentionnée. Les normes ou autres documents normatifs désignés sont utilisés pour le jugement de ce produit.

IT                  Dichiariamo che i prodotti da noi fabbricati, a cui si riferisce la presente dichiarazione, sono conformi ai requisiti fondamentali ai sensi delle disposizioni della direttiva summenzionata. Per la valutazione del prodotto sono stati applicati le norme o i documenti normatyvi citati.

CZ                  Prohlašujeme, že se námi vyráběné výrobky, na které se vztahuje toto prohlášení, shodují se Základními požadavky podle předpisů výše uvedené směrnice. K posouzení výrobku byly použity uvedené normy nebo normatyvy.

ES                  Por la presente declaramos que los productos que nosotros fabricamos, y a los que hace referencia esta declaración, cumplen los requisitos básicos de conformidad con las disposiciones de la directiva arriba mencionada. Las normas o documentos normativos mencionados se aplican en la evaluación de dichos productos.

Biberach

Įsigaliojimo data                    2010-10-02  
 Galiojimo pabaigos data        2017-10-01

CE 0124

DEKRA

Certification GmbH  
 Handwerkstrasse 15  
 D-70565 Štuttgartas

Pardavimo vadybininkas  
 Laimonas Zenkus

/parašas/

Norbert Glaeske  
 Kokybės vadovas

**Tomas Kiškauskas**  
 Med. įrangos inžinierius

Aš, vertėja *Manija Leukulyte*, esu susipažinusi su LR BK 235 straipsniu, kuris numato baudžiamąją atsakomybę už neteisingą vertimą.  
 Vertė..... *[Signature]*

EC-Declaration of Conformity based on 93/42/EEC / Annex II  
EG-Konformitätserklärung basierend auf 93/42/EEG / Anhang II  
CE-Déclaration de conformité basée sur 93/42/EEC / Annexe II  
Dichiarazione di conformità secondo le norme CE 93/42/EEC / Annex II  
Prohlášení o shodě s normami ES založena na 93/42 EEC / Dodatek II  
Declaración de Conformidad de la CE basada en 93/42/EEC / Anexo II  
EK-megfelelőségi nyilatkozat a 93/42/EEC szabványnak megfelelően/II.Függelék



KaVo. Dental Excellence.

**Manufacturer:**

**Kaltenbach & Voigt GmbH  
Bismarckring 39  
D-88400 Biberach**

**MASTERtorque LUX M9000 L  
MASTERtorque LUX M9000 LS  
MASTERtorque LUX M9000 L COLOR (anthracite)  
MASTERtorque LUX M9000 L COLOR (chocolate brown)**

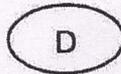
in connection with **MULTiflex (LUX / LED) Coupling**

**Type: 453 N / 454 N / 465 RN / 456 B / 457 / 460 E / 460 LE / 460 LED / 465 LED / 465 LRN / 456 CN / 456 LMN / 456 AN / Midwest**

**93 / 42 / EEC      Medical Devices / Annex I  
EN 1639 : 2009    Medical Devices / Annex IX - Class IIa (Rule 9)  
Dentistry - Instruments**



We declare in sole responsibility that the products manufactured by us to which this statement refers, conform to the essential requirements according to the above-mentioned directive. The specified standards or normative documents were applied to evaluate the product.



Wir erklären in alleiniger Verantwortung, dass die von uns hergestellten Produkte auf die sich diese Erklärung bezieht, mit den grundlegenden Anforderungen gemäß den Bestimmungen der obigen Richtlinie übereinstimmen. Zur Beurteilung des Erzeugnisses wurden die benannten Normen oder normative Dokumente angewandt.



Nous déclarons sous notre seule responsabilité que les produits fabriqués par nos soins, auxquels se réfère cette déclaration, sont conformes aux exigences essentielles selon les dispositions de la directive susmentionnée. Les normes ou autres documents normatifs désignés sont utilisés pour l'évaluation de ce produit.



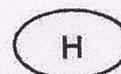
Dichiariamo sotto nostra esclusiva responsabilità che i prodotti da noi fabbricati, a cui si riferisce la presente dichiarazione, sono conformi ai requisiti fondamentali ai sensi delle disposizioni della direttiva summenzionata. Per la valutazione del prodotto sono stati applicati le norme o i documenti normativi citati.



Prohlašujeme s výhradní odpovědností, že se námi vyráběné výrobky, na které se vztahuje toto prohlášení, shodují se základními požadavky podle předpisů výše uvedené směrnice. K posouzení výrobku byly použity uvedené normy nebo normativy.



Por la presente declaramos que en nuestra propia responsabilidad los productos que nosotros fabricamos, y a los que hace referencia esta declaración, cumplen los requisitos básicos de conformidad con las disposiciones de la directiva arriba mencionada. Las normas o documentos normativos mencionados se aplican en la evaluación de dichos productos.



Felelőségünk teljes tudatában nyilatkozatjuk, hogy termékeink, melyekre ez a megfelelőségi nyilatkozat vonatkozik, megfelelnek a fenn említett előírásoknak. Termékeink a fenn említett előírások és szabványok alapján lettek elbírálva.

effective date    Biberach, 02.10.2015  
expiry date      Biberach, 01.10.2018

**Stefan Trampler**  
Director Regulatory Affairs

Pardavimo įrašymas  
Laimonas Žentelis



**Notified Body - Dekra Certification GmbH - Handwerkstrasse 15 - D-70565 Stuttgart - ID-Number: 0124**

90035393 / 503 / 04

EC-Atitikties deklaracija 93/42/EEC / Priedas II  
 EG-Konformitätserklärung basierend auf 93/42/EEG / Anhang II  
 CE-Déclaration de conformité basée sur 93/42/EEC / Annexe II  
 Dichiarazione di conformità secondo le norme CE 93/42/EEC / Annex II  
 Prohlášení o shodě s normami ES založena na 93/42 EEC / Dodatek II  
 Declaración de Conformidad de la CE basada en 93/42/EEC / Anexo II  
 EK-megfelelőségi nyilatkozat a 93/42/EEC szabványnak megfelelően/II.Függelék



KaVo. Dental Excellence.

**Gamintojas:**

**Kaltenbach & Voigt GmbH**  
**Bismarckring 39**  
**D-88400 Biberach**

**MASTERtorque LUX M9000 L**  
**MASTERtorque LUX M9000 LS**  
**MASTERtorque LUX M9000 L COLOR (Atracitos spalvos)**  
**MASTERtorque LUX M9000 L COLOR (šokoladinė ruda)**

jungiamo su **MULTiflex (LUX / LED) jungtimis**

**Type: 453 N / 454 N / 465 RN / 456 B / 457 / 460 E / 460 LE / 460 LED / 465 LED / 465 LRN / 456 CN / 456 LMN / 456 AN / Midwest**

**93 / 42 / EEC Medicinos prietaisai / I priedas**  
**EN 1639 : 2009 Medicinos prietaisai / IX priedas • Ila klasė (9 taisyklė) Stomatologijos-instrumentai**

G

Deklaruojame, kad mūsų gaminami produktai, minimi šiame dokumente atitinka esminius nurodymus direktyvos reikalavimus. Vertinant produktų pasitelkta direktyvoje nurodytais standartais ar normatyviniais aktais.

D

Wir erklären in alleiniger Verantwortung, dass die von uns hergestellten Produkte auf die sich diese Erklärung bezieht, mit den grundlegenden Anforderungen gemäß den Bestimmungen der obigen Richtlinie übereinstimmen. Zur Beurteilung des Erzeugnisses wurden die benannten Normen oder normative Dokumente angewandt.

FR

Nous déclarons sous notre seule responsabilité que les produits fabriqués par nos soins, auxquels se réfère cette déclaration, sont conformes aux exigences essentielles selon les dispositions de la directive susmentionnée. Les normes ou autres documents normatifs désignés sont utilisés pour l'évaluation de ce produit.

IT

Dichiaro sotto nostra esclusiva responsabilità che i prodotti da noi fabbricati, a cui si riferisce la presente dichiarazione, sono conformi ai requisiti fondamentali ai sensi delle disposizioni della direttiva summenzionata. Per la valutazione del prodotto sono stati applicati le norme o i documenti normativi citati.

CZ

Prohlašujeme s výhradní odpovědností, že se námi vyráběné výrobky, na které se vztahuje toto prohlášení, shodují se základními požadavky podle předpisů výše uvedené směrnice. K posouzení výrobku byly použity uvedené normy nebo normativy.

ES

Por la presente declaramos que en nuestra propia responsabilidad los productos que nosotros fabricamos, y a los que hace referencia esta declaración, cumplen los requisitos básicos de conformidad con las disposiciones de la directiva arriba mencionada. Las normas o documentos normativos mencionados se aplican en la evaluación de dichos productos.

H

Felelőségünk teljes tudatában kinyilatkoztatjuk, hogy termékeink, melyekre ez a megfelelőségi nyilatkozat vonatkozik, megfelelnek a fenn említett előírásoknak. Termékeink a fenn említett előírások és szabványok alapján lettek elbírálva.

Isigaliojimo data Biberach, 02.10.2015  
 expiry date Biberach, 01.10.2018  
 Galioja iki Biberach, 01.10.2018

Pardavimo vadybininkas  
 Laimonas Zenkus

**Stefan Trampler**  
 Director Regulatory Affairs

Kopija tikra



EC-Declaration of Conformity based on 93/42/EEC / Annex II  
EG-Konformitätserklärung basierend auf 93/42/EWG / Anhang II  
CE-Déclaration de conformité basée sur 93/42/EEC / Annexe II  
Dichiarazione di conformità secondo le norme CE 93/42/EEC / Annex II  
Prohlášení o shodě s normami ES založena na 93/42 EEC / Dodatek II  
Declaración de Conformidad de la CE basada en 93/42/EEC / Anexo II  
EK-megfelelőségi nyilatkozat a 93/42/EEC szabványnak megfelelően/II.Függelék



KaVo. Dental Excellence.

**Manufacturer:**

**Kaltenbach & Voigt GmbH  
Bismarckring 39  
D-88400 Biberach**

**KaVo PROPHYflex Perio Powder**

**93 / 42 / EEC**

**Medical Devices / Annex I**

**Medical Devices / Annex IX - Class IIa (Rule 9)**

**GB**

We declare in sole responsibility that the products manufactured by us to which this statement refers, conform to the essential requirements according to the above-mentioned directive. The specified standards or normative documents were applied to evaluate the product.

**D**

Wir erklären in alleiniger Verantwortung, dass die von uns hergestellten Produkte auf die sich diese Erklärung bezieht, mit den grundlegenden Anforderungen gemäß den Bestimmungen der obigen Richtlinie übereinstimmen. Zur Beurteilung des Erzeugnisses wurden die benannten Normen oder normative Dokumente angewandt.

**FR**

Nous déclarons sous notre seule responsabilité que les produits fabriqués par nos soins, auxquels se réfère cette déclaration, sont conformes aux exigences essentielles selon les dispositions de la directive susmentionnée. Les normes ou autres documents normatifs désignés sont utilisés pour l'évaluation de ce produit.

**IT**

Dichiariamo sotto nostra esclusiva responsabilità che i prodotti da noi fabbricati, a cui si riferisce la presente dichiarazione, sono conformi ai requisiti fondamentali ai sensi delle disposizioni della direttiva summenzionata. Per la valutazione del prodotto sono stati applicati le norme o i documenti normativi citati.

**CZ**

Prohlašujeme s výhradní odpovědností, že se námi vyráběné výrobky, na které se vztahuje toto prohlášení, shodují se základními požadavky podle předpisů výše uvedené směrnice. K posouzení výrobku byly použity uvedené normy nebo normativy.

**ES**

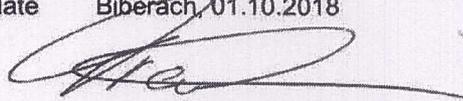
Por la presente declaramos que en nuestra propia responsabilidad los productos que nosotros fabricamos, y a los que hace referencia esta declaración, cumplen los requisitos básicos de conformidad con las disposiciones de la directiva arriba mencionada. Las normas o documentos normativos mencionados se aplican en la evaluación de dichos productos.

**H**

Felelőségünk teljes tudatában nyilatkozatjuk, hogy termékeink, melyekre ez a megfelelőségi nyilatkozat vonatkozik, megfelelnek a fenn említett előírásoknak. Termékeink a fenn említett előírások és szabványok alapján lettek elbírálva.

effective date Biberach, 02.10.2015

expiry date Biberach, 01.10.2018

  
**Stefan Trampler**  
Director Regulatory Affairs



**Notified Body - Dekra Certification GmbH - Handwerkstrasse 15 - D-70565 Stuttgart - ID-Number: 0124**

## Vertimas iš anglų kalbos

EC-Atitikties deklaracija 93/42/EEC I Priedas II  
EG-Konformitätserklärung basierend auf 93/42/EWG / Anhang II  
CE-Déclaration de conformité basée sur 93/42/EEC / Annexe II  
Dichiarazione di conformità secondo le norme CE 93/42/EEC / Annex II  
Prohlášení o shodě s normami ES založena na 93/42 EEC / Dodatek II  
Declaración de Conformidad de la CE basada en 93/42/EEC / Anexo II  
EK-megfelelőségi nyilatkozat a 93/42/EEC szabványnak megfelelően/II.Függelék



KaVo. Dental Excellence.

### Gamintojas:

**Kaltenbach & Voigt GmbH  
Bismarckring 39  
D-88400 Biberach**

## **KaVo PROPHYflex Perio Powder**

**93 / 42 / EEC**

**Medicinos prietaisai / I priedas  
Medicinos prietaisai / IX priedas. klase IIa  
(dalis 9) Odontologijos instrumentai**

GB

Patvirtiname, kad šioje deklaracijoje aprašyti mūsų gaminami gaminiai atitinka esminius aukščiau minėtus direktyvos reikalavimus. Produkto įvertinimui buvo taikomi atitinkami standartai ir norminiai dokumentai.

D

Wir erklären in alleiniger Verantwortung, dass die von uns hergestellten Produkte auf die sich diese Erklärung bezieht, mit den grundlegenden Anforderungen gemäß den Bestimmungen der obigen Richtlinie übereinstimmen. Zur Beurteilung des Erzeugnisses wurden die benannten Normen oder normative Dokumente angewandt.

FR

Nous déclarons sous notre seule responsabilité que les produits fabriqués par nos soins, auxquels se réfère cette déclaration, sont conformes aux exigences essentielles selon les dispositions de la directive susmentionnée. Les normes ou autres documents normatifs désignés sont utilisés pour l'évaluation de ce produit.

IT

Dichiaro sotto nostra esclusiva responsabilità che i prodotti da noi fabbricati, a cui si riferisce la presente dichiarazione, sono conformi ai requisiti fondamentali ai sensi delle disposizioni della direttiva summenzionata. Per la valutazione del prodotto sono stati applicati le norme o i documenti normativi citati.

CZ

Prohlašujeme s výhradní odpovědností, že se námi vyráběné výrobky, na které se vztahuje toto prohlášení, shodují se základními požadavky podle předpisů výše uvedené směrnice. K posouzení výrobku byly použity uvedené normy nebo normativy.

ES

Por la presente declaramos que en nuestra propia responsabilidad los productos que nosotros fabricamos, y a los que hace referencia esta declaración, cumplen los requisitos básicos de conformidad con las disposiciones de la directiva arriba mencionada. Las normas o documentos normativos mencionados se aplican en la evaluación de dichos productos.

H

Felelőségünk teljes tudatában nyilatkozthatjuk, hogy termékeink, melyekre ez a megfelelőségi nyilatkozat vonatkozik, megfelelnek a fenn említett előírásoknak. Termékeink a fenn említett előírások és szabványok alapján lettek elbírálva.

Įsigaliojimo data: 2015.10.02  
Galiojimo pabaigos data: 2018.10.01

Parašas

**Stefan Trampler**  
Direktorius Regulatory Affairs



**Notified Body - Dekra Certification GmbH - Handwerkstrasse 15 - D-70565 Stuttgart - ID-Number: 0124**