

UAB Intersurgical
Amioniu g. 60, Pabrade, LT-18170, Lithuania
T: +370 387 66611 F: +370 387 66622
info@intersurgical.lt www.intersurgical.lt

2018-07-23
No. SI18-279

To whom it may concern

We are pleased to confirm that *Echipamed - Plus S.R.L.* is an official distributor for all Intersurgical product range in Moldova.

Should you have any questions, you may contact me at +370 387 66612.

Sincerely,

Meilutė Milišauskienė
European sales manager
Intersurgical UAB



Sales administrator Renata Pacar, phone: (+00)370 38735113, e-mail: RP@intersurgical.com

EC Certificate Full Quality Assurance System: Certificate GB19/964232

The management system of

Intersurgical Ltd.

Crane House, Molly Millars Lane, Wokingham, Berkshire, RG41 2RZ, UK

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

The scope of registration appears on page 2 of this certificate.

This certificate is valid from 31 October 2019 until 26 November 2022 and remains valid subject to satisfactory surveillance audits.

Issue 1. Certified since 11 January 1995 and first approved by SGS Belgium on 31 October 2019.

This is a multi-site certification. Additional site details are listed on subsequent pages

Certification is based on reports numbered GB/PC 04303

Authorised by

Pieter Weterings
Certification Manager

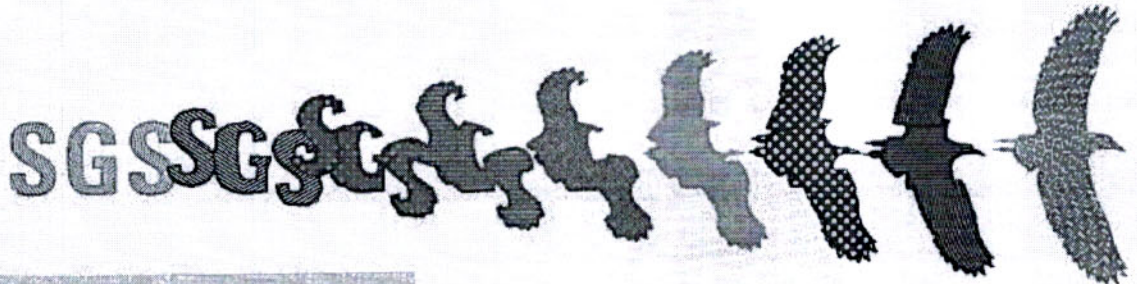


SGS Belgium NV, Notified Body 1639

SGS House Noorderlaan 87 2030 Antwerp Belgium
t +32 (0)3 545-48-48 f +32 (0)3 545-48-49 www.sgs.com

LPM05007 - Certificate CE1639 Annex II-4, EN rev. 02

Page 1 of 2



This document is issued by the Company subject to its General Conditions of Certification Services, unless otherwise agreed, accessible at www.sgs.com/en/ma_and_conditions.htm. Attention is drawn to the limitations of liability, indemnification and jurisdictional issues established therein. The authenticity of this document may be verified at <https://www.sgs.com/en/certified-client-and-product-directory>. Any unauthorized alteration, forgery or falsification of the content or appearance of this document is unlawful and offenders may be prosecuted to the fullest extent of the law.

Intersurgical Ltd.

Directive 93/42/EEC

on medical devices, Annex II (excluding Section 4).

Issue 1

Detailed scope

- Sterile and non-sterile medical devices for respiratory care, in the areas of airway management, anaesthesia, critical care, oxygen and aerosol therapy:**
- Sterile and non-sterile Anaesthetic Breathing Systems
 - Non-sterile Aerosol and Oxygen Masks • Non-sterile Anaesthetic Masks
 - Sterile Bougies and Stylets • Sterile and non-sterile Breathing Systems
 - Non-sterile Bubble Humidifier • Non-sterile Caps
 - Sterile and non-sterile Catheter Mounts
 - Sterile and non-sterile Connectors
 - Sterile and non-sterile Respiratory Filters
 - Non-sterile Flextube Flexible Tubing
 - Non-sterile High Concentration Oxygen Masks • Sterile and non-sterile HME Filters
 - Sterile and non-sterile Humidification Chambers • Sterile I-gel Supraglottic Airways
 - Sterile Laryngeal Mask Airways • Non-sterile Monitoring Lines
 - Non-sterile Nasal Cannulae • Non-sterile Nebulisers
 - Non-sterile Suction and Irrigation Toothbrush
 - Non-sterile Oxygen Lines
 - Non-sterile Repeated Use Breathing Systems
 - Non-sterile Reservoir Bags
 - Non-sterile Manual Resuscitation Systems
 - Non-sterile CO2 Absorbents and Anaesthesia Systems containing Absorbents
 - Non-sterile Suction Tubes and Wands
 - Sterile and non-sterile Suction System • Sterile Endotracheal Tubes
 - Non-sterile Venturi Valves • Non-sterile Wall Nebuliser
 - Non-sterile Water Traps • Non-sterile CPAP and NIV Masks
 - Non-sterile Pressure Limiting Valves • Non-sterile Peep Valves
 - Non-sterile Exhalation Valves
 - Non-sterile One Way Directional Valves
 - Non-sterile nFlow Infant Nasal CPAP device
 - Non-sterile Oxygen Recovery Kits
 - Non-sterile electrically powered moisture condenser for breathing systems
 - Non-sterile Molar Bite Block
 - Non-sterile Mainstream Monitoring (CO2 Cuvette)

Class I sterile: Sterility aspects only - Restricted to the aspects of manufacture concerned with securing and maintaining sterile conditions

- Sterile Guedel Airways

Additional facilities

UAB Intersurgical, Amioniu 60, LT-18170 Pabradė, Lithuania
 Circuit House, Pitronnerie Road, Industrial Estate, St Peter Port,
 Guernsey, GY1 2RL, UK



Certificate GB95/4313



The management system of

Intersurgical Ltd

Crane House, Molly Millars Lane, Wokingham, RG41 2RZ, UK

Unit 1, The Business Centre, Molly Millars Lane,
Wokingham, Berkshire, RG41 2RZ, UK

Brook House, Molly Millars Bridge, Wokingham, Berkshire, RG41 2WY, UK

has been assessed and certified as meeting the requirements of



ISO 9001:2015

For the following activities

The design, manufacture, and supply of medical respiratory products.

This certificate is valid from 08 August 2018 until 08 August 2021 and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 24 July 2021
Issue 26. Certified since 11 January 1995



Authorised by

SGS United Kingdom Ltd
Rossmore Business Park Ellesmere Port Cheshire CH65 3EN UK
t +44 (0)151 350-6666 f +44 (0)151 350-6600 www.sgs.com

HC SGS 9001 2015 0118

Page 1 of 1



This document is issued by the Company subject to its General Conditions of Certification Services accessible at www.sgs.com/terms_and_conditions.htm. Attention is drawn to the limitations of liability, indemnification and jurisdictional issues established therein. The authority of this document may be verified at <http://www.sgs.com/certified-clients-and-products/certified-clients-directory>. Any unauthorized alteration, forgery or falsification of the content or appearance of this document is unlawful and offenders may be prosecuted to the fullest extent of the law.



Certificate GB95/6635

The management system of

Intersurgical Ltd.

Crane House, Molly Millars Lane, Wokingham, Berkshire, RG41 2RZ, UK

has been assessed and certified as meeting the requirements of

ISO 13485:2016 EN ISO 13485:2016

For the following activities

The scope of registration appears on page 2 of this certificate.

This certificate is valid from 19 March 2019 until 26 November 2020
and remains valid subject to satisfactory surveillance audits.

Re certification audit due before 26 November 2019

Issue 29. Certified since 11 January 1995

This is a multi-site certification.

Additional site details are listed on the subsequent page.

Authorised by



SGS United Kingdom Ltd
Rossmore Business Park, Ellesmere Port, Cheshire, CH65 3EN, UK
t +44 (0)151 350-6666 f +44 (0)151 350-6600 www.sgs.com

HC SGS 13485 2016 0118 M2

Page 1 of 2



This document is issued by the Company subject to its General Conditions of Certification Services accessible at www.sgs.com/terms_and_conditions.htm. Attention is drawn to the limitations of liability, indemnification and jurisdictional issues established therein. The authenticity of this document may be verified at <http://www.sgs.com/en/certified-clients-and-products/certified-client-directory>. Any unauthorized alteration, forgery or falsification of the content or appearance of this document is unlawful and offenders may be prosecuted to the fullest extent of the law.

Certificate GB95/6635, continued



Intersurgical Ltd.

ISO 13485:2016 EN ISO 13485:2016

Issue 29

Detailed scope

Design and manufacture of sterile and non-sterile medical devices for respiratory care, in the areas of airway management, anaesthesia, critical care and oxygen and aerosol therapy.

Additional facilities

**Intersurgical Ltd.
Circuit House, Pitronnerie Road, Industrial Estate,
St Peter Port, Guernsey, GY1 2RL, UK**

**UAB Intersurgical
Armonijų g. 60, LT-18170 Pabradė, Lithuania**



0005

