



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

No. Issued To: CE 69386 CooperSurgical Inc., also trading as Ackrad Laboratories, Prism Healthcare, Milex, Medscand, Wallach Surgical Devices, SAGE In-Vitro Fertilization and Lone Star Medical Products 95 Corporate Drive, Trumbull, Connecticut 06611 USA

In respect of:

See certificate scope page.

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 2797):

Gary C Stade

Gary E Slack, Senior Vice President Medical Devices

First Issued: 2002-08-30

Date: 2020-03-13

Expiry Date: 2024-05-26

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Certificate No: CE 69386

Certificate Scope:

The manufacture of Ophthalmic, Gynaecology and dermatology Cryosurgery Systems and accessories.

The manufacture of sterile and non-sterile obstetrics and gynaecology devices including;

- •Uterine Manipulators/Injectors and their accessories including colpotomizer cups
- •Vaginal Pessaries
- Amniocentesis Needles
- •Vacuum assisted Delivery systems and accessories
- •Infant warmers
- •Fetal and Vascular Ultrasound Dopplers

The manufacture of sterile and non-sterile Artificial Reproductive Technology (ART) devices including;

- •Oocyte recovery needles
- Incubators
- Micropipettes
- Stripper Tips

The manufacture of sterile general surgery devices including;

- •Esophageal ballon Catheters
- •Laparoscopic cannulas/trocars
- •Laparoscopic Port Closure Systems
- •Stays, retractors and retractor kits

Those aspects of Annex V relating to securing and maintaining sterility in the manufacture of;

Intrauterine Insemination Catheters, Embryo Replacement Catheters, Pasteur pipettes, Assisted Hatching pipettes, Laparoscopic Smoke Evacuation Systems, Hysteroscopy Saline/Air Contrast Injectors, Diagnostic Surveillance Cannulas, Adapter Drapes, Hysterosalpingography Catheters, Bi-nasal Oxygen Cannulas, Cervical Dilators, Gynaecological tissue sampling devices.

Those aspects of Annex V relating to metrology in the manufacture of; Vaginal Pessary Fitting Kit and Contraceptive Diaphragm fitting sets

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Supplementary Information to CE 69386

Issued To:

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Number	Device Name	Intended purpose per IFU
Class IIa		
MD 1103	Medical Heat Packs	
MD 1302	Ultrasound dopplers	
MD 1104	Cryosurgical systems	
MD 1402	IVF Incubators G185 & G210	
MD 0106	Adult Esophageal Balloon Catheter Set	1
MD 0106	Milex® Pessaries (Short term)	
MD 0106	Uterine Manipulators/Injectors and Accessories	
MD 0106	Fetal vacuum extraction system and Accessories	
MD 0106	Carter-Thomason CloseSure System®	
MD 0106	Stays and Retractors	
MD 0106	Apple-Hunt Secondary Trocar Cannula	
MD 0106	Carter-Thomason® II Port Closure System Suture Passer	
MD 0102	Wallace® Amniocentesis Needle	
MD 0109	Wallace® Oocyte Recovery Set	
MD 0109	Micropipettes	L CDDE
MD 0109	Stripper Tips	

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Number	Device Name	Intended purpose per IFU	
Class Is		NOW THE REACT	
MD 0106	Lone Star Retractor Rings		
MD 0109	Pasteur Pipettes		
MD 0109	Origio Assisted Hatching Pipettes		
MD 0101	INCA® Nasal Cannula Set with sterile nasal prongs		
MD 0109	Intrauterine Insemination Catheters		
MD 0106	Cytology Sampling devices		
MD 0106	Endometrial Suction Curette	((()))	
MD 0106	Hysterosalpingography Catheter		
MD 0106	Laparoscopic Smoke Evacuation Systems	0	
MD 0106	Wallach Disposable OS Finders		
MD 0106	CSI OS Finders	J 1	
MD 0106	ABBI (Air Bubble Base Infuser)		
MD 0106	EndoSee® Cannula		
MD 0106	Sterile Adapter Drapes		
MD 0109	Embryo Replacement Catheters	S- ESSE	
Class Im			
MD 0107	Vaginal Diaphragms Fitting Sets		
MD 0106	Milex Pessaries Fitting Kit		
Circt Techody	2002_09_20 Data: 2020_02_12	Evoin Data: 2021-05-26	

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

CE 69386

List of Significant Subcontractors

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

Certificate No: Date:

Issued To:

2020-03-13 CooperSurgical Inc., also trading as Ackrad Laboratories, Prism Healthcare, Milex, Medscand, Wallach Surgical Devices, SAGE In-Vitro Fertilization and Lone Star Medical Products 95 Corporate Drive, Trumbull, Connecticut 06611 USA

Subcontractor:

Service(s) supplied

Manufacture

Cooper Medical SRL Edificio N° B49, 51 Ave 0 Parque Industrial Zona Franca Coyol La Garita, Alajuela Costa Rica

CooperSurgical Distribution B.V Celsiusweg 35 5928 PR VenIo The Netherlands

Drummond Scientific 500 Parkway Broomall PA 19008 USA

Emergo Europe Prinsessegracht 20 2514 AP The Hague The Netherlands **EU Representative**

Crucial Supplier

EU Representative

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Subcontractor:

Isomedix Operation, Inc. North Facility 1880 Industrial Drive Illinois 60048 USA

Isomedix Operations, Inc 3459 South Clinton Avenue South Plainfield New Jersey 07080 USA Service(s) supplied Gamma Irradiation

Gamma Sterilization

ETO Sterilization

Isomedix Operations, Inc. 435 Whitney Street Northborough Massachusetts 01532 USA Gamma Irradiation Gamma Sterilization

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USA

List of Significant Subcontractors

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Subcontractor:

Service(s) supplied

Isomedix Operations, Inc. 9 Apollo Drive Whippany New Jersey 07981 USA

Lone Star Medical Products 11211 Cash Road Stafford Texas 77477 USA

Origio a/s Knardrupvej 2 2760 Måløv Denmark Gamma Irradiation Gamma Sterilization

Manufacture

EU Representative

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Subcontractor:

Service(s) supplied

Pristech Products Inc. 6952 Fairgrounds Parkway San Antonio Texas 78238 USA

Research Instruments Ltd. Bickland Industrial Park Falmouth TR11 4TA United Kingdom

Sterigenics US,LLC 84 Park Road Queensbury NY 12804 USA Manufacture

Manufacture

ETO Sterilization

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Subcontractor:

Service(s) supplied

ETO Sterilization

Sterilization Services of Tennessee, Inc. 2396 Florida Street Memphis Tennessee 38109 USA

UNISIS CORP. Saitama Plant 2675-1 Nishikata Koshigaya-shi Saitama 343-0822 Japan **Crucial Supplier**

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Certificate No: Date:

CE 69386

2020-03-13

Issued To:

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Date	Reference Number	Action
30 August 2002	4316414	Original issue to CooperSurgical Inc for the manufacture of Fetal Dopplers and Sterile Disposable Uterine Manipulator-Injectors.
04 October 2002	4386116	Amended certificate to add Sterile Colpotomizer Systems to scope.
12 November 2002	4386116	Amended certificate to add the trading name of Ackrad Laboratories Inc. Also amended scope to include INCA Nasal Cannulae Set with sterile nasal prongs and Sterile Bronchitrac Suction Catheters, Esophageal Balloon Catheters, Elliptosphere Catheters, Tampa Catheters and H/S Catheter sets with Kraton Balloons.
28 January 2004	4487082	Amended certificate to add the trading name of Prism Healthcare. Also amended scope to include Gelpacks for heat therapy, sterile and non-sterile Vacuum Assisted Delivery Pumps, sterile Vacuum Assisted Delivery Extractor Cups and Delivery Kits. Also added a new subcontractor Pristech Products Inc as manufacture of gelpacks for heat therapy.
25 February 2004	n/a	Correction of typographical error to address.

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Certificate No: CE 69386 Date: 2020-03-13 Issued To: CooperSurgical Inc., also trading as Ackrad Laboratories, Prism Healthcare, Milex, Medscand, Wallach Surgical Devices, SAGE In-Vitro Fertilization and Lone Star Medical Products 95 Corporate Drive, Trumbull, Connecticut 06611 USA

Date	Reference Number	Action
22 July 2004	4593904	Amended certificate to add the trading name of SAGE In-Vitro Fertilization Inc. Also, amended scope to include sterile oil and sterile water for Tissue Culture for use in in-vitro fertilization, and Sterile Phosphate Buffered Saline (PBS) for use in in-vitro fertilization Also, modified the Supplementary Information to highlight Class I Sterile devices.
27 September 2004	4634606	Amended certificate to include Vascular Dopplers within the scope.
30 November 2004	4647383	Amended certificate to include Ophthalmic Cryosurgery Systems within the scope.
27 April 2005	4657997	Amended certificate to include trading name of Milex and amended scope to include Sterile Curettes and Contraceptive Diaphragm fitting sets.
19 September 2005	4739488	Extension of scope to include Fiberoptic Cystometry System, Sterile Fiberoptic Catheter and Cytology Sampling devices.
16 July 2007	7073121	Certificate renewal.
		Extension to scope to include Embryo Transfer Catheters.
30 October 2007	7110759	Extension to scope to include Intrauterine Insemination Catheters.

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Certificate No: CE 69386 Date: 2020-03-13 Issued To: CooperSurgi also trading Prism Health Wallach Sur SAGE In-Vite

2020-03-13 CooperSurgical Inc., also trading as Ackrad Laboratories, Prism Healthcare, Milex, Medscand, Wallach Surgical Devices, SAGE In-Vitro Fertilization and Lone Star Medical Products 95 Corporate Drive, Trumbull, Connecticut 06611 USA

Date	Reference Number	Action
23 May 2008	7213846	Extension to scope to include Cryosurgery Systems for gynaecology and dermatology. Also addition of Wallach Surgical Devices as a trading name.
23 June 2008	7226858	Extension to scope to include CT CloseSure Systems and CT CloseSure Systems XL.
22 October 2008	7284056	Extension to scope to include endometrial suction curettes.
31 March 2010	7507745	Addition of Lone Star Medical Products as a trading name and addition of Sterile Surgery Kits, Retractor Kits, Stays, Dilamezinsert and Sterile Retractors to scope. These were covered by Lone Star Medical Products BSI CE 553016 issued on 12 October 2009 which was originally transferred from SGS where Lone Star Medical Products, Inc. was certified since 4 February 1998. Steris Isomedix Services, Inc., Coventry, RI - Sterilization was removed and three new Subcontractors were added: Steris Isomedix, So. Plainfield, NJ - ETO Sterilization; Leisegang Feinmechanik GmbH, Berlin, Germany - EU Representative; and Lone Star Medical Products, Stafford, TX – Manufacture. With this revised certificate, CE 69386, the previous certificate CE 553016 is cancelled.

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EC Certificate - Production Quality Assurance Certificate History

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Date	Reference Number	Action
02 August 2010	7548521	Addition of Sterigenics US, LLC, 84 Park Road, Glen Falls, New York, 12804, to the list of significant subcontractors for ETO sterilization.
24 February 2011	7650994	Addition of Laparoscopic Smoke Evacuation Systems to scope. Addition of significant subcontractor Biotest Laboratories, Inc.
13 February 2012	7752683	 Revised format of supplementary pages for clarity. Addition of vaginal pessaries to scope. Revision of the following devices to "Those aspects of Annex V relating to securing and maintaining sterility in the manufacture of". INCA Nasal Cannulae Set with sterile nasal prongs. Intrauterine Insemination Catheters. Revision of the following devices to "Manufacture of". Vacuum Assisted Delivery Pumps. Sterile Elliptosphere Catheters. Sterile Tampa Catheters. Sterile H/S Catheters sets with Kraton Balloons.

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Date	Reference Number	Action
13 February 2012	7752683	- Removal from certificate scope of.
(continued)		Oil for Tissue Culture for use in in-vitro fertilization.
		Phosphate Buffered Saline (PBS) for use in in-vitro fertilization.
		Water for Tissue Culture for use in in-vitro fertilization.
		- Addition of Steris Isomedix, Libertyville as a significant subcontractor.
		Removal of Steris Isomedix, Waukegan and Isomedix Operation Inc as significant subcontractors.
		Removal of sterilisation and testing activities for significant subcontractor Biotest Laboratories.
		Clarification of sterilisation method (Gamma Irradiation) for significant subcontractor Steris Isomedix, Northborough.
12 June 2012	7828419	Extension to scope of manufacturing to include Trocars, Cannulas and Trocar/Cannula Kits and Retractors.
		Addition of 'Emergo Europe, Netherlands' as EU Representative.

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Date	Reference Number	Action
09 August 2012	7868380	Certificate Renewal.
		Removal of 'Biotest Laboratories, Inc' as significant subcontractor.
23 July 2013	8022947	Addition of Vaginal Pessary Fitting Kit.
29 November 2013	8075064	Revision of the following devices to "Those aspects of Annex V relating to securing and maintaining sterility in the manufacture of".
		Elliptosphere catheters, <i>H/S</i> catheter sets with kraton balloons and tampa catheters.
		Removal of Sterile Bronchitrac suction catheters from scope.
10 June 2015	7982958	Addition of Limb Plethysmography Systems.
08 September 2015	8413506	Addition of Sterile ABBI.
06 November 2015	8429989	Addition of Endosee Cannulas.
11 March 2016	8485008	Removal of SAGE In-Vitro Fertilization as significant subcontractor.
23 June 2016	8548511	Addition of Sterile Os Finder Cervical Dilator, Sterile Disposable Smoke Evacuation Tubing with Attached Speculum Tubing, and Sterile Adapter Drape.

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Date	Reference Number	Action
07 August 2017	8695527	Certificate Renewal. Change of address for Emergo Europe. Change of address for Sterigencics US, LLC. Change of name for Steris Isomedix Services to Isomedix Operations, Inc.
		Removal of Sterile Disposable Smoke Evacuation Tubing with Attached Speculum Tubing.
26 January 2018	8887080	Addition of IVF Incubators. Addition of Research Instruments as a significant subcontractor.
23 October 2018	8922778	Addition of Amniocentesis Needles, Embryo Replacement Catheters, and Oocyte Recovery Needles.
		Addition of Cooper Medical SRL, Sterilization Services of Tennessee Inc., and UNISIS CORP as significant subcontractors.
19 February 2019	9719699	Traceable to NB 0086.
24 January 2020	3143385	Removal of Leisegang Feinmechanik GmbH EU Representative. Addition of CooperSurgical Distribution B.V. as EU Representative.

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	USA

Date	Reference Number	Action
Current	9770590	Certificate renewal.
		Origio a/s added as EU Representative.
		Drummond Scientific added as Crucial Supplier
		Isomedix Operations, Inc. (9 Apollo Drive) added as significant subcontractor for Gamma Irradiation/Gamma Sterilization.
		Removal of Sterile Surgery Kits, Dilamezinsert, Limb Plethysmography Systems, Fiberoptic Cystometry Systems and Fiberoptic Catheters from scope.
		Embryo Transfer Catheters removed from scope as it is covered by Embryo replacement catheters.
		Added Micropipettes, Stripper Tips and Pasteur/Assisted Hatching pipettes to scope.
		The scope has been reformatted to better reflect the certified devices and make the expression more understandable.

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