

## 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 00493

Date:

2020-03-15

Issued To:

3M Company

3M Health Care

dba 3M Consumer Health Care

2510 Conway Ave.

Saint Paul Minnesota 55144 USA

Subconti actor	Su	bcon	tractor:
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Service(s) supplied

3M Company 905 Adams St. SE Hutchinson Minnesota 55350 USA Manufacture

3M Deutschland GmbH Healthcare Business Carl-Schurz - Str. 1 41453 Neuss Germany Control of Sterilization EU Representative Secondary Packaging

3M Deutschland GmbH Health Care Business Plant Kamen Edisonstraβe 6 59174 Kamen Germany Manufacture

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#### Subcontractor:

Service(s) supplied

3M Company 10351 West 70th Street Eden Prairie Minnesota 55344 USA

Manufacture

3M Company 5400 Paris Road Columbia Missouri 65202 USA

Manufacture

3M Company 601 22nd Ave. South **Brookings** South Dakota 57006 USA

**Gamma Sterilization** Manufacture

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#### Supplementary Information to CE 00493

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NBOG code(s)	Device description	
Class Is		
MDS 7006	3M <sup>™</sup> Cavilon <sup>™</sup> No Sting Barrier Film (Wipes/wands)	
MDS 7006	3M™ Bair Hugger™ Warming Blankets (Sterile)	
MDS 7006	3M™ Coban™ Self-Adherent Wrap (Sterile)	
MDS 7006	3M™ Steri-Strip™ Closures	
MDS 7006	3M™ Medipore™ +Pad Cloth Adhesive Wound Dressing	
MDS 7006	3M <sup>™</sup> Steri-Drape <sup>™</sup> Surgical Drape and Accessories	
MDS 7006	3M™ Skin Stapler Remover	
MDS 7006	3M™ Nexcare™ Bandages	

First Issued: 1995-02-01 Date: 2020-03-15 Expiry Date: 2024-05-26

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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.



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NBOG code(s)	Device description	
Class IIa		
MD 0301	3M™ Tegaderm™ Transparent Film Dressing	
MD 0301	3M™ Tegaderm™ I.V. Advanced Securement Dressing and 3M™ Tegaderm™ Diamond Pattern Film Dressing	
MD 0301	3M™ Tegaderm™ HP Transparent Film Dressing Frame Style	
MD 0301	3M™ Tegaderm™ Transparent Film Dressing First-Aid Style	
MD 0301	3M™ Tegaderm™ I.V. Transparent Film Dressing with Adhesive-Free Window	
MD 1301	3M™ Littmann® Electronic Stethoscope and software	
MD 0302	3M <sup>™</sup> Precise <sup>™</sup> Disposable Skin Stapler with Staples	
MD 0303	3M™ Steri-Drape™ Isolation Bag and Wound Edge Protector Drape	
MD 0303	3M™ Cavilon™ Advanced Skin Protectant	
MD 0301	3M™ Tegaderm™ + Pad Film Dressing with Nonadherent Pad	
MD 0108	3M™ Curos™ Disinfecting Caps, Tips, and Cap Strips	

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Information and Contact; BSI, Say Building, John M. Keynesplein 9, 1066 EP Amsterdam, The Netherlands Tel: + 31 20 346 0780 BSI Group The Netherlands B.V. registered in The Netherlands under 33264284.

A member of BSI Group of Companies.



Certificate No: CE 00493

### Certificate Scope:

The manufacture of sterile transparent wound dressings with and without pads, electronic stethoscopes and associated software, sterile skin staplers and sterile drapes (wound protector, isolation bag), barrier film dressings, and sterile disinfecting port protector devices.

Those aspects of Annex V relating to securing and maintaining sterility in the manufacture of wound closures, wound dressings, drapes, barrier film wraps, staple removers, patient warming blankets and securement devices.

Those aspects of manufacturing relating to obtaining sterility in the assembly of procedure packs in accordance with Article 12 of the Medical Device Directive.

First Issued: 1995-02-01 Date: 2020-03-15 Expiry Date: 2024-05-26

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No.

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Issued To:

3M Company 3M Health Care

dba 3M Consumer Health Care

2510 Conway Ave. Saint Paul

Minnesota 55144 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 E Slack, Senior Vice President Medical Devices

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Date: 2020-03-15

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