



Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

No. CE 664635

Issued To: **Terumo Medical Corporation**

265 Davidson Avenue, Suite 320

Somerset New Jersey 08873 USA

In respect of:

Design, development and manufacture of sterile vascular closure devices

on the basis of our examination of the quality assurance system under the requirements of Council Directive 93/42/EEC, Annex II excluding section 4. The quality assurance system meets the requirements of the directive. For the placing on the market of class III products an Annex II section 4 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

First Issued: **2017-01-20** Date: **2019-07-18** Expiry Date: **2022-01-19**

...making excellence a habit."

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.

Gay C Stade





Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

List of Significant Subcontractors

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

Certificate No: **CE 664635**

Date: **2019-07-18**

Issued To: **Terumo Medical Corporation**

265 Davidson Avenue, Suite 320

Somerset New Jersey 08873 USA

Subcontractor:

Service(s) supplied

DSM Biomedical 735 Pennsylvania Drive

Exton PA 19341 USA **Animal Tissues / Derivatives**

St. Jude Medical 14901 DeVeau Place Minnetonka Minnesota 55345-2126 USA Manufacture

St. Jude Medical Costa Rica Ltda. Edificio #44, Calle 0, Ave. 2 Zona Franca El Coyol, Alajuela Costa Rica **Manufacture**

...making excellence a habit."





Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

List of Significant Subcontractors

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

Certificate No: **CE 664635** 2019-07-18

Issued To: **Terumo Medical Corporation**

265 Davidson Avenue, Suite 320

Somerset **New Jersey** 08873 USA

Subcontractor:

Date:

Service(s) supplied

Sterigenics US, LLC 1700 College Boulevard West Memphis AR 72301 **USA**

Gamma Sterilization

Sterigenics US, LLC 1003 Lakeside Drive Gurnee

Illinois 60031 **USA**

Gamma Sterilization

Synergy Health AST SRL B13.1 Street 4, Avenue 1 El Coyol Free Zone 20102 El Coyol Alajuela Costa Rica

ETO Sterilization

...making excellence a habit."





Directive 93/42/EEC on Medical Devices, Annex II excluding Section 4

List of Significant Subcontractors

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

Certificate No: **CE 664635**Date: **2019-07-18**

Issued To: **Terumo Medical Corporation**

265 Davidson Avenue, Suite 320

Somerset New Jersey 08873 USA

Subcontractor:

Service(s) supplied

Terumo Europe, N.V. Interleuvenlann 40, B-3001

Leuven Belgium **EU Representative**

Terumo Puerto Rico LLC Innovation Street Lot 21 Caguas West Industrial Park Caguas 00725 Puerto Rico **Manufacture**

...making excellence a habit."





EC Certificate - Full Quality Assurance System **Certificate History**

Certificate No:

CE 664635

Date:

2019-07-18

Issued To:

Terumo Medical Corporation 265 Davidson Avenue, Suite 320

Somerset

New Jersev

08873

USA

Date	Reference Number	Action
20 January 2017	8645594	First issue.
20 October 2017	8794520	Update name and address of subcontractor Terumo Puerto Rico LLC.
27 February 2019	8798156	Traceable to NB 0086.
Current	9659326	Change in legal manufacturer address Updated address of Terumo Puerto Rico LLC to align with BSI ISO certificates.

...making excellence a habit.™ 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.