



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

No. Issued To: CE 554734 Terumo Corporation 44-1, 2-chome Hatagaya Shibuya-ku Tokyo 151-0072 Japan

In respect of:

The Design and Manufacture of Balloon Dilatation Catheters, PTCA Guidewires, Angiographic Catheters, MicroGuide catheters, Coronary Imaging Catheters and coronary optical coherence tomography system.

Those aspects of Annex II related to securing and maintaining the sterility of the MDU cover, Extension Wires, and related accessories.

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 C Stade

Gary E Slack, Senior Vice President Medical Devices

First Issued: 2009-10-30

Date: 2019-08-12

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.





Supplementary Information to CE 554734

Issued To:

Terumo Corporation 44-1, 2-chome Hatagaya Shibuya-ku Tokyo 151-0072 Japan

Number	Device Name	Intended purpose per IFU
Class III		
	RyujinPlus	See CE 554735
	Tazuna	See CE 554735
	Hiryu	See CE 599214
	RyujinPlus OTW	See CE 578316
	Accuforce	See CE 608484
	Ryurei	See CE 661655
	Progreat	See CE 580672
	Finecross MG	See CE 597867
	Runthrough NS	See CE 613749
	FastView	See CE 585621

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Issued To:

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Number	Device Name	Intended purpose per IFU
Class IIa		
MD 1202	LUNAWAVE	
Class Is		
MD 0106	RunthroughNS Extension wire	
MD 0106	Fast View MDU cover	

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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: Date:

Issued To:

2019-08-12 Terumo Corporation 44-1, 2-chome Hatagaya Shibuya-ku Tokyo 151-0072 Japan

CE 554734

Subcontractor:

Service(s) supplied

SUZUKI Co., Ltd. 2150-1 Ogawara Suzaka-shi Nagano 382-8588 Japan

Terumo Corporation Ashitaka Plant 150, Maimaigi-cho, Fujinomiya City, Shizuoka Prefecture 418-0015 Japan

Terumo Europe N.V. Interleuvenlaan 40 3001 Leuven Belgium Manufacture

Design Development ETO Sterilization Manufacture

EU Representative

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List of Significant Subcontractors

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

Certificate No: Date:

Issued To:

2019-08-12 Terumo Corporation 44-1, 2-chome Hatagaya Shibuya-ku Tokyo 151-0072 Japan

CE 554734

Subcontractor:

Ueda Japan Radio Co., Ltd. 2805-72 Nagase Ueda-shi Nagano 386-0407 Japan Service(s) supplied

Manufacture

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EC Certificate - Full Quality Assurance System Certificate History

Certificate No: Date:

Issued To:

CE 554734 2019-08-12 Terumo Corporation 44-1, 2-chome Hatagaya Shibuya-ku Tokyo 151-0072

Japan

Date	Reference Number	Action
30 October 2009	7443727	First Issue – Transfer from another Notified Body.
17 September 2010	7560390	Certificate renewal.
23 December 2011	7778290	Addition of "Angiographic Catheters" to the scope of the certificate. Additional service supplied for ETO sterilization at the Terumo Ashitaka Plant.
30 March 2012	7730762	Update to scope of certificate to add Coronary Imaging Catheters.
21 December 2012	7916383	Extension to scope to include LUNAWAVE.
18 April 2013	7948395 7959985	Optical Coherence Tomography System (LUNAWAVE) was introduced under 7916383 in Dec 2012. Brand name 'LUNAWAVE' has now been removed from scope. This does not affect the device types covered by the certificate.
		Extension of scope to include Class I sterile MDU cover and accessories.
4 June 2013	7974363	Extension to scope to include micro-guide catheters.
4 June 2014	8164373	Certificate renewal.
1 August 2014	8196034	Addition of "PTCA Guidewires" and "sterility ofExtension Wires" to the scope.

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EC Certificate - Full Quality Assurance System Certificate History

Certificate No:	CE 554734
Date:	2019-08-12
Issued To:	Terumo Corporation 44-1, 2-chome Hatagaya Shibuya-ku Tokyo 151-0072 Japan

Date	Reference Number	Action
27 April 2018	8942575	Added design and development service to Terumo Ashika Plant subcontractor.
04 March 2019	7778938	Traceable to NB 0086.
Current	9789827	Certificate Renewal. Added products table and subcontractors Ueda Japan Radio and SUZUKI.

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