



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

No. CE 554734

Issued To: **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 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.

Gary C Stade





Supplementary Information to CE 554734

Issued To: **Terumo Corporation**

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

| 100 (100 m) (100 m) | | | | |
|---------------------|----------------|--------------------------|--|--|
| 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 | | |

First Issued: **2009-10-30** Date: **2019-08-12** Expiry Date: **2024-05-26**

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Supplementary Information to CE 554734

Issued To: Terumo Corporation

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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 | 500 |

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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: **CE 554734**Date: **2019-08-12**

Issued To: **Terumo Corporation**

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

Subcontractor:

Service(s) supplied

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

Terumo Corporation Ashitaka Plant 150, Maimaigi-cho, Fujino

150, Maimaigi-cho, Fujinomiya City, Shizuoka Prefecture 418-0015

Japan

Design
Development
ETO Sterilization
Manufacture

Terumo Europe N.V. Interleuvenlaan 40 3001 Leuven Belgium **EU Representative**

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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: CE 554734 Date: 2019-08-12

Issued To: **Terumo Corporation**

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

Subcontractor:

Service(s) supplied

Ueda Japan Radio Co., Ltd.

2805-72 Nagase

Ueda-shi

Nagano

386-0407 Japan

Manufacture

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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 |
|-------------------|---------------------|---|
| 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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