

EC Certificate

Directive 93/42/EEC Annex II, excluding Section 4 **Full Quality Assurance System Medical Devices**

Registration No.: HD 60149405 0001

Report No.:

12018179 053

Manufacturer:

OLYMPUS MEDICAL SYSTEMS CORP.

2951 Ishikawa-cho. Hachioji-shi, Tokyo 192-8507 Japan

Products:

Design and Development, Manufacture of Medical Endoscopy

Systems, Diagnostic, Operation and Treatment Products

(see attachments for products included)

Replaces Approval, Registration No.: HD 60144066 0001

Expiry Date:

2024-05-26

The Notified Body hereby declares that the requirements of Annex II, excluding section 4 of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex II, section 5 of the aforementioned directive. For placing on the market of class III devices covered by this certificate an EC design-examination certificate according to Annex II, section 4 is required.

Effective Date:

0/020 d 04 08 S TÜV, TUEV and TUV are registered trademarks. Unisation and application requires prior approva

2020-05-12

Date:

2020-05-12

Notified Body TÜVRheinland enifizierungs6

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 90431 Nürnberg TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.



TÜV Rheinland LGA Products GmbH Tillystraße 2, 90431 Nürnberg

Doc. 1/1, Rev.0

Attachment to Certificate

Registration No.:

HD 60149405 0001

Report No.:

12018179 053

Manufacturer:

OLYMPUS MEDICAL SYSTEMS CORP.

2951 Ishikawa-cho, Hachioji-shi, Tokyo 192-8507 Japan

Products included:

Medical Endoscopy Systems:

- -Endoscopes
- -Endotherapy Devices
- -Imaging Processors
- -Pumps for Endoscopy
- -Light Sources
- -Position Detecting Units
- -Electrothermal Cautery Units
- -Integrated Endosurgery Systems
- -Endoscopic Regulation/Control Units

Electrosurgical Equipment

Probes and Transducers for Ultrasonic Lithotriptors

Laparoscopic Insufflators

Ultrasound Surgical Equipment

Ultrasonic Surgical System generator

Ultrasonic Surgical System transducer

Hard-tissue ultrasonic surgical system holder/tip

Disinfecting Units

Date: 2020-05-12

Capsule Endoscopes and Systems

Ultrasound Diagnostic Imaging Systems

Notified Body

M.Sc. M. Aihara

⊤ÜVRheinland



EC Certificate

Directive 93/42/EEC Annex II, excluding Section 4 **Full Quality Assurance System Medical Devices**

Registration No.: HD 60151129 0001

Report No.:

21232416 020

Manufacturer:

Olympus Winter & Ibe GmbH

Kuehnstr. 61 22045 Hamburg Deutschland

Products:

medical endoscopy, surgical, diagnostic, and treatment

systems

(see attachment for products and sites included)

Replaces Approval, Registration No.: HD 60104211 0001

Expiry Date:

2024-05-26

The Notified Body hereby declares that the requirements of Annex II, excluding section 4 of the directive 93/42/EEC have been met for the listed products. The above named manufacturer has established and applies a quality assurance system, which is subject to periodic surveillance, defined by Annex II, section 5 of the aforementioned directive. For placing on the market of class III devices covered by this certificate an EC design-examination certificate according to Annex II, section 4 is required.

Effective Date:

2020-08-12

Date:

2020-08-12

Roland Aruber

TÜVRheinland

Notified Body

TÜV Rheinland LGA Products GmbH - Tillystraße 2 - 9043 🖟 TÜV Rheinland LGA Products GmbH is a Notified Body according to Directive 93/42/EEC concerning medical devices with the identification number 0197.



Doc. 1/3, Rev.0

TÜV Rheinland LGA Products GmbH Tillystraße 2, 90431 Nürnberg

Attachment to Certificate

Registration No.:

HD 60151129 0001

Report No.:

21232416 020

Manufacturer:

Olympus Winter & Ibe GmbH

Kuehnstr. 61 22045 Hamburg Deutschland

Products included:

- Telescopes, Videoscopes and Fiberscopes
- Electrosurgical Instruments
- Electrosurgical Forceps
- Electrosurgical Hand Pieces
- Electrosurgical Units
- Electrosurgical Electrodes
- Electrosurgical Cables and Adapters, Active Cords
- Peristaltic Pump Units
- Peristaltic Tubing Sets
- Invasive Access Devices (which would encompass obturators, sheaths, trocars, bridges and irrigation ports)
- Endoscopy Instruments
 (which would encompass cannulas, injection needles, ballon dilators, adaptors, tubes and non-active instruments)

Date: 2020-08-12

91.V 1

Notified Body

Roland Gruber

TÜVRheinlar



TÜV Rheinland LGA Products GmbH Tillystraße 2, 90431 Nürnberg Doc. 2/3, Rev.0

Attachment to Certificate

Registration No.:

HD 60151129 0001

Report No.:

21232416 020

Manufacturer:

Olympus Winter & Ibe GmbH

Kuehnstr. 61 22045 Hamburg Deutschland

Products included:

- Working Elements and Inserts
- Bladder Syringes
- Washer-Disinfectors
- Adaptors for Washer-Disinfectors
- Fluid Management Systems used in Endoscopy
- Light Sources

For the following medical devices the scope covers only the aspects of manufacture concerned with securing and maintaining sterile conditions:

- sterile Caps

Date: 2020-08-12

Roland Gruber

Notified Body





TÜV Rheinland LGA Products GmbH Tillystraße 2, 90431 Nürnberg

Doc. 3/3, Rev.0

Attachment to Certificate

Registration No.:

HD 60151129 0001

Report No.:

21232416 020

Manufacturer:

Olympus Winter & Ibe GmbH

Kuehnstr. 61 22045 Hamburg Deutschland

Sites included:

Olympus Winter & Ibe GmbH Rheinstr. 8 14513 Teltow Germany

- Design and development, manufacturing of medical devices for electrosurgery and for endoscopic applications

Date: 2020-08-12

Gruber

Notified Body

Certificate

Quality Management System EN ISO 13485:2016

Registration No.:

SX 2158445-1

Organization:

OLYMPUS MEDICAL SYSTEMS CORP.

2951 Ishikawa-cho Hachioji-shi, Tokyo 192-8507 Japan

Scope:

Design and Development, Manufacture, Distribution, Service, Quality Assurance, Regulatory Affairs, Planning, Delivery support of Endoscopes, Endotherapy devices, Light Sources, Imaging Processors, Endoscope Position Detecting Units, Electrothermal Cautery Units, Integrated Endosurgery

Systems, Endoscopic Regulation/Control Units, Camera

Heads/Pumps/Monitors/Recorders for Endoscopy, Electrosurgical Equipment, Capsule Endoscopes and Systems, Laparoscopic Insufflators, Ultrasound Diagnostic Imaging Systems, Disinfecting Units, Ultrasound Surgical Equipment, Ultrasonic surgical system generator, Ultrasonic surgical system transducer, Hard-tissue ultrasonic surgical system holder/tip, Probes and Transducers for Ultrasonic Lithotriptors, Sterile Non Active Instruments used in conjunction with Endoscopes, Sterile Endotherapy Devices used in conjunction with Endoscopes, Sterile Non Active Devices used in conjunction with Medical Ultrasound Diagnostic Imaging Systems and Water Container, Water Supply

Tube, Water Feeding valve and Foot Switch for Pump

The Certification Body of TÜV Rheinland LGA Products GmbH certifies that the organization has established and applies a quality management system for medical devices.

Proof has been furnished that the requirements specified in the abovementioned standard are fulfilled. The quality management system is subject to yearly surveillance.

Tüvnheinland

Report No.:

150242490-301

Effective date:

2022-03-07

Expiry date:

2024-07-26

Issue date:

2022-03-07



Ning N. C. Chang TÜV Rheinland LGA Products GmbH Marierungsster illystraße 2 · 90431 Nürnberg · Germany

TÜVRheinland

Certificate

Quality Management System EN ISO 13485:2016

Registration No.:

SX 2158445-1

Organization:

OLYMPUS MEDICAL SYSTEMS CORP.

2951 Ishikawa-cho Hachioji-shi, Tokyo 192-8507 Japan

The scope of certification also covers the following:

| No. | Facility | Scope |
|-----|--|---|
| /01 | c/o OLYMPUS MEDICAL SYSTEMS CORP. Ishikawa Office 2951 Ishikawa-cho, Hachioji-shi, Tokyo 192-8507 Japan | Design and Development, Service, Administration, Management, Quality Assurance, Regulatory Affairs, Planning, Delivery support |
| /02 | c/o OLYMPUS MEDICAL SYSTEMS CORP. Shinjuku Monolith Office Shinjuku Monolith, 3-1 Nishishinjuku 2-chome, Shinjuku-ku, Tokyo 163-0914 Japan | Quality Assurance for domestic market |
| /03 | c/o OLYMPUS MEDICAL SYSTEMS CORP. Shirakawa Facility 3-1 Okamiyama, Odakura, Nishigo- mura, Nishishirakawa-gun, Fukushima 961-8061 Japan | Service and Quality Assurance |

Report No.:

150242490-301

Effective date:

2022-03-07

Expiry date:

2024-07-26

Issue date:

2022-03-07



TÜVRheinland LGA Products GmbH
TÜVstraße 2 · 90431 Nürnberg · Germany



Certificate

Quality Management System EN ISO 13485:2016

Registration No.:

SX 2158445-1

Organization:

OLYMPUS MEDICAL SYSTEMS CORP.

2951 Ishikawa-cho Hachioji-shi, Tokyo 192-8507 Japan

The scope of certification also covers the following:

/04

c/o OLYMPUS MEDICAL SYSTEMS

Quality Assurance

CORP. Aizu Facility 3-1-1 Niiderakita,

Aizuwakamatsu-shi, Fukushima

965-8520 Japan

Report No.:

150242490-301

Effective date:

2022-03-07

Expiry date:

2024-07-26

Issue date:

2022-03-07



TÜVRheintand

TÜV Rheinland LGA Products GmbH

Tillystraße 2 · 90431 Nürnberg · Germany



Certificate

Quality Management System EN ISO 13485:2016

Registration No.:

SX 2158445-1

Organization:

OLYMPUS MEDICAL SYSTEMS CORP.

2951 Ishikawa-cho Hachioji-shi, Tokyo 192-8507 Japan

The scope of certification also covers the following:

/05

c/o OLYMPUS MEDICAL SYSTEMS

CORP.

Aomori Facility 2-248-1 Okkonoki, Kuroishi-shi, Aomori 036-0357 Japan Quality Assurance of Endoscopes, Endotherapy devices, Light Sources, Imaging Processors, Endoscope Position Detecting Units, Electrothermal Cautery Units, Integrated Endosurgery Systems, Endoscopic Regulation /Control Units, Camera Heads/Pumps/Monitors /Recorders for Endoscopy, Electrosurgical Equipment, Capsule Endoscopes and Systems, Laparoscopic Insufflators, Ultrasound Diagnostic Imaging Systems, Disinfecting Units and Ultrasound Surgical Equipment, Probes and Transducers for Ultrasonic Lithotriptors, Sterile Non Active Instruments used in conjunction with Endoscopes, Sterile Endotherapy Devices used in conjunction with Endoscopes, Sterile Non Active Devices used in conjunction with Medical Ultrasound Diagnostic Imaging Systems and Water Container, Water Supply Tube, Water Feeding valve and Foot Switch for Pump

Design and Development of Endotherapy Devices, Sterile Endotherapy Devices used in conjunction with Endoscopes

Report No.:

150242490-301

Effective date:

2022-03-07

Expiry date:

2024-07-26

Issue date:

2022-03-07



TÜVRheinland

TÜVRheinland

TÜVRheinland

TÜV Rheinland LGA Products GmbH

Tillystraße 2 · 90431 Nürnberg · Germany