



FUJIFILM Corporation
26-30, NISHIAZABU 2-CHOME, MINATO-KU,
TOKYO 106-8620 JAPAN

To Whom It May Concern

Date: November 14, 2013

CERTIFICATE

We, FUJIFILM Corporation, hereby certify that the attached "EC Certificate" is a copy of original certificate.

*Certificate Registration Number : G1 12 09 20011 029

With best regards,

FUJIFILM Corporation



Yasuhiro Aihara
Senior Manager
Endoscopy Systems Div.
Medical Systems Business Div.



Product Service

EC Certificate

Full Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 12 09 20011 029

Manufacturer: FUJIFILM Corporation
26-30, Nishiazabu 2-Chome
Minato-Ku, Tokyo
106-8620 JAPAN

EC-Representative: FUJIFILM Europe GmbH
Heesenstr. 31
40549 Düsseldorf
GERMANY

Product Category(ies): Computed Radiography Console,
Diagnostic X-ray Equipment,
Digital Mammography System and
related equipment, Software for Diagnostic
imaging Workstations, Ultrasound Diagnostic
Imaging Equipment, Endoscopes for medical use
and their related equipment and accessories,
Ultrasound Endoscopes and their related
equipment and accessories

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.: TAQ235005952A

Valid from: 2012-11-22

Valid until: 2017-11-21

Date, 2012-11-15

Hans-Heiner Junker



TÜV SÜD Product Service GmbH Is Notified Body with identification no. 0123

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Product Service

EC Certificate**Full Quality Assurance System**

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 12 09 20011 029**Facility(ies):**

FUJIFILM Corporation
26-30, Nishiazabu 2-Chome, Minato-Ku, Tokyo,
106-8620 JAPAN

FUJIFILM Corporation, Medical Systems Research
& Development Center
798, Miyanodai, Kaisei-Machi, Ashigarakami-Gun,
Kanagawa, 258-8538 JAPAN



Registered No. 1399

NOTARIAL CERTIFICATE

This is to certify that KENICHI KAMOI, an agent of YASUHIRO AIHARA, Senior Manager, Endoscopy Systems Div., Medical Systems Business Div. of FUJIFILM CORPORATION, has stated in my very presence that said YASUHIRO AIHARA has acknowledged himself to have signed and sealed the attached document.

Dated this 15th day of November, 2013.



KO ITO

Notary

1-18-1 Shimbashi, Minato-ku, Tokyo, Japan
Tokyo Legal Affairs Bureau

添付書面の作成者である 富士フイルム株式会社 メディカルシステム事業部 内視鏡システム部 統括マネージャー 相原康宏 の代理人 鴨井謙一 は本職に対し、前記 相原康宏 がその署名捺印を自認している旨、陳述した。

よって、これを認証する。

平成25年 11 月 15 日、本公証人役場において
東京都港区新橋1丁目18番1号
東京法務局所属

公 証 人
Notary

伊藤 剛
KO ITO

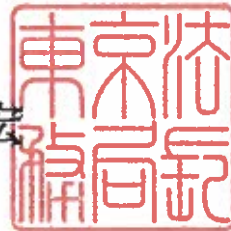
証 明

上記署名は、東京法務局所属公証人の署名に相違ないものであり、かつ、その押印は、真実のものであることを証明する。

平成25年 11 月 15 日

東京法務局長

石田一宏



APOSTILLE

(Convention de La Haye du 5 octobre 1961)

1. Country: JAPAN

This public document

2. has been signed by KO ITO

3. acting in the capacity of Notary of the Tokyo Legal Affairs Bureau

4. bears the seal/stamp of KO ITO, NOTARY

Certified

5. at Tokyo

6. 11/15/2013

7. by the Ministry of Foreign Affairs

8. 13-**N0029781**

9. Seal/stamp:

10. Signature



A. Ogawa

Ayako OGAWA

For the Minister for Foreign Affairs