FUJIFILM

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

Y. Aihara

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

Digitally signed by Botnaru Andrei
Date: 2019.03.18 13:44:27 EET
Reason: MoldSign Signature
Location: Moldova

ZERTIFIKAT ◆ CERTIFICATE ◆ 認証書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICA

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

499915

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

TÜV®

ZERTIFIKAT ◆ CERTIFICATE ◆ 認証証書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICA

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.

TOKYO
LEGAL AFFAIRS BUREAU
NOTARY
1-18-1 SHIMBASHI MINATO-KU
TOKYO JAPAN

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

平成２５年登簿第 1390号

認　　証

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 岡

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-№029781
9. Seal/stamp:
10. Signature

MINISTRY OF FOREIGN AFFAIRS JAPAN

A. Ogawa

Ayako OGAWA

For the Minister for Foreign Affairs