EC CERTIFICATE

umber: 76910CE01

Production Quality Assurance

Directive 93/42/EEC on Medical devices, Annex V
(Devices in Class IIa, IIb or III)

Manufacturer:

FUJIFILM Corporation

26-30 Nishiazabu
2-Chome Minato-Ku
Tokyo 106-8620
Japan

For the product category(ies)

X-ray films

DEKRA grants the right to use the EC Notified Body Identification Number illustrated below to accompany the CE Marking of Conformity on the products concerned conforming to the required Technical Documentation and meeting the provisions of the EC-Directive which apply to them:

0344

Documents, that form the basis of this certificate:

Certification Notice 76910CN, initially dated 24 December 1997
Addendum, initially dated 15 December 2000

DEKRA hereby declares that the above mentioned manufacturer fulfils the relevant provisions of 'Besluit Medische Hulpmiddelen', the Dutch transposition of the Council Directive 93/42/EEC of June 14, 1993 concerning Medical devices, including all subsequent amendments. The manufacturer has implemented a quality assurance system for the manufacture and final inspection for the above mentioned product category in accordance to the provisions of Annex V of Council Directive 93/42/EEC of June 14, 1993 and is subject to periodical surveillance. For placing on the market of Class III or Class IIb devices an additional EC type-examination certificate according to Annex III is mandatory. The necessary information related to the quality assurance system of the manufacturer, including facilities and the reference to the relevant documentation, of the products concerned and the assessments performed, are stated in the Certification Notice which forms an integrative part of this certificate.

This certificate is valid until: 1 April 2024
Issued for the first time: 24 December 1997
Reissued: 1 April 2019

DEKRA Certification B.V.

B.T.M. Holtus
Managing Director

J.A. van Vugt
Certification Manager

Digitally signed by Antonova Anna
Date: 2021.01.06 16:16:57 EET
Reason: MoldSign Signature
Location: Moldova

DEKRA Certification B.V. is Notified Body with ID no 0344

DEKRA Certification B.V. Meander 1051, 6825 MJ Arnhem P.O. Box 5185, 6802 ED Arnhem, The Netherlands

ADDENDUM

Belonging to certificate: 76910CE01

CE MARKING OF CONFORMITY
MEDICAL DEVICES

X-ray films

Issued to:

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

This certificate covers the following product(s):

X-ray film

Initial date: 15 December 2000
Revision date: 2 October 2006

DEKRA Certification B.V.

B.T.M. Holtus
Managing Director

J.A. van Vugt
Certification Manager

DEKRA Certification B.V. is Notified Body with ID no 0344

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