TÜV SÜD
Product Service

EC Certificate

Production Quality Assurance System

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

No. G2 18 01 20011 045

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):	**Imaging Plates for Computed Radiography**

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

Report No.:	**JAQ235032409**
Valid from:	**2018-05-14**
Valid until:	**2023-05-13**

Digitally signed by Grabazei Alexandru
Date: 2020.10.08 10:35:51 EEST
Reason: MoldSign Signature
Location: Moldova

Date, 2018-04-12

Stefan Preiß

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

ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ СЕРТИФИКАТ ◆ CERTIFICADO ◆ CERTIFICAT

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Facility(ies):

FUJIFILM Corporation
798, Miyanodai, Kaisei-Machi, Ashigarakami-Gun, Kanagawa,
258-8538 JAPAN

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