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

Digitally signed by Botnaru Andrei
Date: 2022.11.29 12:41:04 EET
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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

Date, 2018-04-12

Stefan Preiß

04052767797903

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

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Facility(ies):

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

TÜV SÜD Product Service GmbH · Zertifizierstelle · Ridlerstraße 65 · 80339 München · Germany

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