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TÜV SÜD Product Service

EC Certificate

Production Quality Assurance System
Directive 93/42/EEC on Medical Devices, Annex V
(Devices in class I with measuring function)

No. G2M 020011 0049 Rev. 00

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):	**Workstation, Computed Radiography Console**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for the manufacture in accordance with MDD Annex V. This quality assurance system covers those aspects of manufacture concerned with the metrological requirements of the respective devices / device categories and conforms to the requirements of this Directive. It is subject to periodical surveillance. See also notes overleaf.

Report No.:	**JAQ235032836**
Valid from:	**2018-08-28**
Valid until:	**2023-02-28**

Date, **2018-08-28**

I. Preiß

Stefan Preiß

Digitally signed by Grabazei Alexandru
Date: 2020.10.08 10:36:15 EEST
Reason: MoldSign Signature
Location: Moldova

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

-/-

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