



Certificate

No. Q5 057385 0018 Rev. 00

Holder of Certificate: FUJIFILM TECHNO PRODUCTS CO., LTD.
1250, Takematsu
Minamiashigara-shi, Kanagawa
250-0111 JAPAN

Certification Mark:



Scope of Certificate: Production and Distribution of Clinical Chemistry Analyzer, Computed Radiographic Equipment (including CR Console), Workstations, Medical Imaging Equipment, Ultrasound Diagnostic Imaging Equipment, IP Cassettes for X-ray Storage Media (Imaging Plate), Medical Imaging Plate, Digital Radiographic Equipment, Peripherals of Medical Endoscope, and Digital Mammography and Accessories

The Certification Body of TÜV SÜD Product Service GmbH certifies that the company mentioned above has established and is maintaining a quality management system, which meets the requirements of the listed standard(s). See also notes overleaf.

Report No.: JAQ235034823

Valid from: 2018-11-20
Valid until: 2020-11-30

Date, 2018-11-20

Stefan Preiß

TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD TÜV SÜD
 ZERTIFIKAT ◆ CERTIFICATE ◆ 認證證書 ◆ CERTIFICADO ◆ CERTIFIKAT ◆ CERTIFICATE

Certificate

No. Q5 057385 0018 Rev. 00

Applied Standard(s): EN ISO 13485:2016
Medical devices - Quality management systems -
Requirements for regulatory purposes
(ISO 13485:2016)
DIN EN ISO 13485:2016

Facility(ies): FUJIFILM TECHNO PRODUCTS CO., LTD.
Takematsu Factory
1250, Takematsu, Minamiashigara-shi, Kanagawa,
250-0111 JAPAN

FUJIFILM TECHNO PRODUCTS CO., LTD. Tohoku
Factory Hanamaki Site
2-1-3, Kitayuguchi, Hanamaki-Shi, Iwate, 025-0301
JAPAN

-/-