





Certificate

No. Q5 043398 0289 Rev. 00

Holder of Certificate:

Nipro Corporation

NIPRO

3-9-3, Honjo-Nishi, Kita-ku

Osaka 531-8510

JAPAN

Facility(ies):

Nipro Corporation Odate Factory

8-7, Hanukiyachi, Niida, Odate-shi, Akita, 018-5794 JAPAN

Certification Mark:



Scope of Certificate:

Design and Development, Production, Sterilization (EO, gamma and e-beam), Packaging, and Distribution of Sterile and Non-sterile Single Use Medical Products for Blood Sampling, Hemodialysis, Infusion, Injection,

Perfusion, Plasmapheresis, Transfusion, Cardiovascular, Vascular, and ART (Assisted

Reproductive Technology),

Caps for the Application of Pharmaceutical Products, In-Vitro Diagnostic Kits for Biochemistry, Immunology

and Genetics, Medical Grade Silicone

Applied Standard(s):

EN ISO 13485:2016

Medical devices - Quality management systems -

Requirements for regulatory purposes

(ISO 13485:2016)

DIN EN ISO 13485:2016

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.:

JNQ235033902

Valid from:

2018-09-01

Valid until:

2021-08-31

Date,

2018-08-28

Stefan Preiß

1. Pumil

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