



Product Service

EC Certificate

Full Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4)
(Devices in Class IIa, IIb or III)

No. G1 17 04 24736 060

Manufacturer:

KANEKA Corporation

3-18, 2-Chome, Nakanoshima, Kita-ku
Osaka-city, OSAKA
530-8288 JAPAN



EC-Representative:

KANEKA PHARMA EUROPE N.V.

Nijverheidsstraat 16
2260 Westerlo-Oevel
BELGIUM

Product

Category(ies):

**Suction Catheter Set, Catheter for Angioplasty and
Balloon Dilation Catheter for Angioplasty**

The Certification Body of TÜV SÜD Product Service GmbH declares that the aforementioned manufacturer has implemented a quality assurance system for design, manufacture and final inspection of the respective devices / device categories in accordance with MDD Annex II. This quality assurance system conforms to the requirements of this Directive and is subject to periodical surveillance. For marketing of class III devices an additional Annex II (4) certificate is mandatory. See also notes overleaf.

Report No.:

JNQ235029570

Valid from:

2017-08-17

Valid until:

2022-01-21

Date, 2017-08-17

Stefan Preiß



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

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Full Quality Assurance System

Directive 93/42/EEC on Medical Devices (MDD), Annex II excluding (4) (Devices in Class IIa, IIb or III)

No. G1 17 04 24736 060

Facility(ies):

KANEKA Corporation
3-18, 2-Chome, Nakanoshima, Kita-ku, Osaka-city, OSAKA,
530-8288 JAPAN

KANEKA Corporation Osaka Plant
5-1-1, Torikai-Nishi, Settsu-city, OSAKA, 566-0072 JAPAN

KANEKA Medix Corporation Kanagawa Plant
225-1, Aza Deguchi, Yamakita, Yamakita-machi,
Ashigara-Kami-gun, KANAGAWA, 258-0113 JAPAN

KANEKA PHARMA VIETNAM CO., LTD.
35 VSIP Street 6, Vietnam - Singapore Industrial Park, An Phu
Ward, Thuan An Town, Binh Duong Province, VIETNAM



Product Service

CERTIFICATE

No. Q5 17 04 24736 065

Holder of Certificate: **KANEKA Corporation**
KANEKA 3-18, 2-Chome, Nakanoshima, Kita-ku
 Osaka-city, OSAKA
 530-8288 JAPAN

Certification Mark:



Scope of Certificate: Design and Development, Production and Distribution of
 Medical Device for Selective Plasma Component
 Adsorption, Medical Device for Blood Purification,
 Neuro Surgical Products, Catheters for
 Interventional Radiology, Surgical Products,
 Silicone Tubing Ophthalmic Products,
 Cell Separation Device
 Design and Development and Distribution of
 Plasma Separator, Blood Tubing Lines, Apheresis Unit
 Distribution of Blood Flowmeter

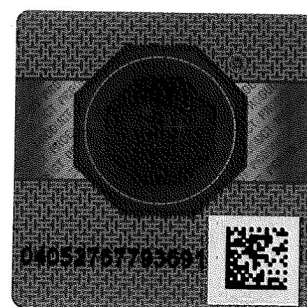
The Certification Body of TÜV SÜD Product Service GmbH certifies that the company mentioned above has established and is maintaining a quality management system, which meets the requirements of the listed standard(s). See also notes overleaf.

Report No.: JNQ235029570

Valid from: 2017-09-01
 Valid until: 2020-08-31

Date, 2017-08-17

S. Preiß
 Stefan Preiß



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DAKKS
 Deutsche
 Akkreditierungsstelle
 D-ZM-11321-01-00



Product Service

CERTIFICATE**No. Q5 17 04 24736 065**

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):

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 3-18, 2-Chome, Nakanoshima, Kita-ku, Osaka-city, OSAKA,
 530-8288 JAPAN

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 225-1, Aza Deguchi, Yamakita, Yamakita-machi,
 Ashigara-Kami-gun, KANAGAWA, 258-0113 JAPAN

KANEKA Medix Corporation Osaka Office
 3-18, 2-Chome, Nakanoshima, Kita-ku, Osaka-city, OSAKA,
 530-8288 JAPAN

KANEKA PHARMA VIETNAM CO., LTD.
 35 VSIP Street 6, Vietnam - Singapore Industrial Park, An Phu
 Ward, Thuan An Town, Binh Duong Province, VIETNAM

KANEKA Corporation Tokyo Office
 12-32, 1-Chome, Akasaka, Minato-ku, Tokyo, 107-6028 JAPAN

KANEKA Medix Corporation Tokyo Office
 12-32, 1-Chome, Akasaka, Minato-ku, Tokyo, 107-6028 JAPAN

KANEKA Medix Corporation Tokyo Logistics Center
 1-4-3, Katsu-shima, Shinagawa-ku, Tokyo, 140-0012 JAPAN



Product Service

EC Certificate

EC Design-Examination Certificate

Directive 93/42/EEC on Medical Devices (MDD), Annex II (4)
(Devices in Class III)

No. G7 18 05 24736 068

Manufacturer:

KANEKA Corporation

3-18, 2-Chome, Nakanoshima, Kita-ku
Osaka-city, OSAKA
530-8288 JAPAN



EC-Representative:

KANEKA PHARMA EUROPE N.V.

Nijverheidsstraat 16
2260 Westerlo-Oevel
BELGIUM

Product:

Catheters for Single Use

The Certification Body of TÜV SÜD Product Service GmbH declares that a design examination has been carried out on the respective devices in accordance with MDD Annex II (4). The design of the devices conforms to the requirements of this Directive. For marketing of these devices an additional Annex II certificate is mandatory. See also notes overleaf.

Report no.:

713121823

Valid from:

2018-06-19

Valid until:

2023-05-21

Date, 2018-06-19

Stefan Preiß



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Product Service

EC Certificate**EC Design-Examination Certificate**

Directive 93/42/EEC on Medical Devices (MDD), Annex II (4)
(Devices in Class III)

No. G7 18 05 24736 068

Model(s):**Dual Lumen Catheters**

- Crusade Catheter KMF0114A

- FineDuo Catheter NC-D724

Parameters:

Effective length 1400 ± 20 mm

Maximum outer diameter 1.07 ± 0.02 mm

For guide wire size 0.014 inch (0.36 mm)

For guide catheter size 0.056 inch (1.44 mm)

Facility(ies):

KANEKA Corporation

3-18, 2-Chome, Nakanoshima, Kita-ku, Osaka-city, OSAKA,
530-8288 JAPAN

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5-1-1, Torikai-Nishi, Settsu-city, OSAKA, 566-0072 JAPAN

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