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Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
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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 067727 0007 Rev. 01

Manufacturer: **Venner Medical (Deutschland) GmbH**
Gut Uhlenhorst 4
24229 Dänischenhagen
GERMANY

Facility(ies): Venner Medical (Deutschland) GmbH
Gut Uhlenhorst 4, 24229 Dänischenhagen, GERMANY

Product Category(ies): Infusion pumps and related disposables

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.: 713163321

Valid from: 2019-08-21

Valid until: 2024-05-26

Date, 2019-08-21

Stefan Preiß
Head of Certification/Notified Body

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