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Zentralstelle der Länder
für Gesundheitsschutz
bei Arzneimitteln und
Medizinprodukten
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ZLG-BS-244.10.08



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 050643 0002 Rev. 01

Manufacturer: **UAB VILTECHMEDA**

Mokslininku 6
08412 Vilnius
LITHUANIA

Facility(ies):

UAB VILTECHMEDA
Mokslininku 6, 08412 Vilnius, LITHUANIA

Product Category(ies): **Infusion syringe pumps, infusion pumps, applied part
(docking station, monitoring - configuration software)**

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.: 713159145

Valid from: 2019-10-01

Valid until: 2024-05-26

Date, 2019-09-13

Stefan Preiß
Head of Certification/Notified Body