



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 078672 0014 Rev. 01

Manufacturer: **Shenzhen Urion Technology Co., Ltd.**
Floor 4-6th of Building D
Jiale Science & Technology Industrial Zone
No.3, ChuangWei Road
Heshuikou Community, MaTian Street
GuangMing New District
518106 Shenzhen
PEOPLE'S REPUBLIC OF CHINA

Product Category(ies): Digital Blood Pressure Monitors, Infrared Thermometer, Compressor Nebulizers

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. All applicable requirements of the testing and certification regulation of TÜV SÜD Group have to be complied with. For details and certificate validity see: www.tuvsud.com/ps-cert?q=cert:G10786720014Rev.01

Report No.: GZ2117902

Valid from: 2021-04-12

Valid until: 2024-05-26

Date, 2021-04-12

Christoph Dicks
Head of Certification/Notified Body