



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)

G1 078672 0012 Rev. 00

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

Shenzhen Urion Technology Co., Ltd. Facility(ies):

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

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.: GZ18179CN01

Valid from: 2018-08-06

Valid until: 2022-02-07

2018-08-06

Date,

1. Pumil

Stefan Preiß