



Certificate

No. Q5 005136 0001 Rev. 02

Holder of Certificate: **Shenzhen Witleaf Medical Electronics Co., Ltd.**

Room 1201, Building 1
Senyang Electronic Technology Park
West Area, Guangming Hi-tech Park
Tianliao Community, Yutang Street
Guangming District
518132 Shenzhen
PEOPLE'S REPUBLIC OF CHINA

Certification Mark:



Scope of Certificate: **Design and Development, Production and Distribution of Patient Monitor, Rapid Intervention Capnograph, Fingertip Pulse Oximeter, Handheld Pulse Oximeter**

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). 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:Q5 005136 0001 Rev. 02

Report No.: GZ2135701

Valid from: 2022-06-01

Valid until: 2025-04-14

Date, 2022-06-01

Christoph Dicks
Head of Certification/Notified Body

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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): Shenzhen Witleaf Medical Electronics Co., Ltd.
Room 1201, Building 1, Senyang Electronic Technology Park,
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See Scope of Certificate