

Certificate

Quality Management System EN ISO 13485:2016 + AC:2018 + A11:2021 ISO 13485:2016

Registration No.: SX 2120274-1

Certificate Holder: Shenzhen Viatom Technology Co., Ltd.
901, Building West, Lepu Tower, No.66
Xingke Road, Xili Community,
Xili Street, Nanshan District,
Shenzhen,
518055 Guangdong
P.R. China

Scope: Design and Development, Manufacture and Distribution of
Vital Signs Monitors, Pulse Oximeters, Blood Pressure
Monitors, Dynamic ECG Recorders, ECG Recorders, Wireless
Dynamic Multi-parameter Holters, Non-invasive Ventilators
and Blood Pressure Monitors with ECG

The Certification Body of TÜV Rheinland LGA Products GmbH certifies that the organization has established and applies a quality management system for medical devices.
Proof has been furnished that the requirements specified in the abovementioned standard are fulfilled. The quality management system is subject to yearly surveillance.

Report No.: 10927775-120

Effective date: 2026-09-02

Expiry date: 2029-09-01

Issue date: 2026-08-21

Replaces certificate SX 2120274-1 issued 2025-12-10

This certificate can be validated on <https://www.certipedia.com>


Joanna Huang
TÜV Rheinland LGA Products GmbH
Tillystraße 2 · 90431 Nürnberg · Germany

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The scope of certification also covers the following sites:

No.	Facility	Scope
/01	c/o Shenzhen Viatom Technology Co., Ltd. 901, Building West, Lepu Tower, No.66 Xingke Road, Xili Community, Xili Street, Nanshan District, Shenzhen, 518055 Guangdong P.R. China	Aspects related to Design and Development and Distribution
/02	c/o Shenzhen Viatom Technology Co., Ltd. Floor 3, Building 9, Baiwangxin Industrial Park, No.1002 Songbai Road, Yangguang Community, Xili Street, Nanshan District, Shenzhen 518055 Guangdong P.R. China	Aspects related to Manufacture

This certificate can be validated on <https://www.certipedia.com>