





Certificate

No. Q5 107512 0001 Rev. 00

Holder of Certificate: ViVest Medical Technology Co., Ltd Unit 205,206,207,B1 Building, SIP BioBay No.218, Xinghu Street Suzhou Industrial Park, Suzhou Area Suzhou Free Trade Pilot Zone 215123 Jiangsu

Facility(ies): ViVest Medical Technology Co., Ltd Unit 205,206,207,B1 Building, SIP BioBay, No.218, Xinghu Street, Suzhou Industrial Park, Suzhou Area, Suzhou Free Trade Pilot Zone, 215123 Jiangsu, PEOPLE'S REPUBLIC OF CHINA

PEOPLE'S REPUBLIC OF CHINA

See Scope of Certificate

EN ISO 13485

Certification Mark:

Scope of Certificate:

Design and Development, Production and Distribution of Automated External Defibrillator

Applied Standard(s):

2021-04-28

EN ISO 13485:2016 Medical devices - Quality management systems -Requirements for regulatory purposes (ISO 13485:2016) DIN EN ISO 13485:2016

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:Q5107512.0001 Rev. 00

Report No.: Valid from: Valid until:

Date,

SH20160801 2021-04-28 2024-04-27

Christoph Dicks Head of Certification/Notified Body