Certificate Number CN19/41019

The quality management system of

Shenzhen Jumper Medical Equipment Co., Ltd.

D Building, No. 71, Xintian Road, Fuyong Street, Baoan,
Shenzhen, Guangdong, 518103, P.R. China

Dun & Bradstreet D-U-N-S identification number: 52-921-7911

has been audited against the criteria stated below and found to conform to those criteria for the scope contained in this certificate

MDSAP(ISO 13485:2016)

Australia:
Therapeutic Goods (Medical Devices) Regulations, 2002, Schedule 3 Part 1 (excluding Part 1.6) – Full Quality Assurance Procedure [including design]
Brazil:
RDC ANVISA n. 16/2013 / RDC ANVISA n. 23/2012 / RDC ANVISA n. 67/2009
Canada:
Medical Devices Regulations – Part 1 SOR 98/282
Japan:
MHLW Ministerial Ordinance 169, Article 4 to Article 68 / PMD Act
United States:
21 CFR 820 / 21 CFR 803 / 21 CFR 806 / 21 CFR 807 – Subparts A to D

For the following activities and devices

Design & Manufacture of Infrared Thermometer, Fingertip Pulse Oximeter, Fetal Doppler, Electronic Blood Pressure Monitor and Electronic Pulse Stimulator

This certificate is valid from
Effective Date: 2019-02-19 until Expiry Date: 2022-02-18
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 2021-09-14
Issue 1. Certified since 2019-02-19

Authorised by J Hall
Business Manager

Jonathan M. Hall

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