EC Certificate Full Quality Assurance System: Certificate CN19/41011

The management system of

Shenzhen Hingmed Medical Instrument Co., Ltd.

4th Floor, Zhonghangfeixiang Building, No.371, Guangshen Road,
Baoan District, Shenzhen, Guangdong, 518102, P.R. China

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex II (excluding Section 4)

For the following products

Ambulatory blood pressure monitor (Model: ABP-03)
Wearable ambulatory blood pressure monitor (Model: WBP-02A)
Clinical Automatic Blood Pressure Monitor
(Model: DBP-01HP, DBP-01P)
Upper Arm Blood Pressure Monitor (Model:Q06, Q06B)

Where the above scope includes class III medical device(s), a valid EC Design Examination Certificate according to Annex II (Section 4) is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 09 April 2021 until 06 August 2023
and remains valid subject to satisfactory surveillance audits.
Issue 3. Certified since since 07 August 2015.

Certification is based on reports numbered CN/SZX 50055

Authorised by

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