EC Certificate Full Quality Assurance System: Certificate CN15/30453

The management system of

Shenzhen Quality Medical Technology Co., Ltd.

5/F, Plant C3, No.1, Nuclear Power Industrial Zone,
Guanlan Shijing Community, Guanlan Street, Longhua New Area,
Shenzhen, Guangdong, 518110, 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

Non Sterile Single use Electrosurgical Neutral Electrode (Model: GP202B-A, GP202M-A, GP202M-P, GP202B-P, GP202M-I, GP202B-I, GP202M-AC, GP202B-AC, GP202M-PC, GP202B-PC, GP202M-IC, GP202B-IC)

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 16 June 2017 until 15 June 2022
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 14 March 2020
Issue 2. Certified since 8 April 2015

Certification is based on reports numbered CN/ZSN 15989

Authorised by

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