

EC Certificate Full Quality Assurance System: Certificate CN14/30268

The management system of

Guangzhou Longest Science & Technology Co., Ltd.

5&6F, Building B4, No.11, Kaiyuan Avenue, Science City,
Guangzhou Hi-tech Industrial Development Zone,
Guangzhou, 510530 Guangzhou, 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

Electrical Extracorporeal Shock Wave Therapy Device (driven by compressed air) for the treatment of chronic muscular pain and tendon disorders (Model: LGT-2500A, LGT-2500AL, LGT-2500B, LGT-2500BL, LGT-2510A, LGT-2510AL, LGT-2510B, LGT-2510BL, LGT-2500S)

Compression Therapy Device LGT-2200WM, LGT-2200WML, LGT-2200S, LGT-2200SL, LGT-2200L and LGT-2200LL intended for:

Prophylaxis of Deep vein thrombosis (DVT) (Post-stroke deep vein thrombosis, Post-traumatic deep vein thrombosis) and Prophylaxis and treatment of edema (Chronic venous edema, Lymphedema, Post-mastectomy lymphedema) and LGT-2200DVT intended for:

Prophylaxis of Deep vein thrombosis (DVT)

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 17 May 2016 until 3 March 2019 and remains valid subject to satisfactory surveillance audits.

Re certification audit due before 15 November 2016

Issue 5. Certified since 3 March 2014

Certification is based on reports numbered CN/CAN 16140

Authorised by

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