

EC Certificate Full Quality Assurance System: Certificate CN19/41109

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, 510530, Guangzhou, Guangdong Province, 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

The scope of registration appears on page 2 of this certificate.

This certificate is valid from 16 December 2019 until 28 January 2024 and remains valid subject to satisfactory surveillance audits.

Issue 1. Certified since 03 March 2014 and first certified by SGS Belgium NV since 16 December 2019.

Certification is based on reports numbered CN/CAN 16140

Authorised by

SGS Belgium NV, Notified Body 1639

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Science & Technology Co., Ltd.

Directive 93/42/EEC

on medical devices, Annex II (excluding Section 4).

Issue 1

Detailed scope

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

Compression Therapy Device LGT-2200WM, LGT-2200WML,
LGT-2200L, LGT-2200LL and LGT-2200SP 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);
LGT-2200DVT and LGT-2201DVT 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.