

The management system of

Sky Medical Technology Limited, also trading as Firstkind Limited

Hawk House, Peregrine Business Park, Gomm Road,
High Wycombe, HP13 7DL, UK

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

Portable electro-stimulation devices: to increase blood circulation, for the prevention of venous thrombosis, for the prevention and treatment of oedema, for promoting wound healing, for the treatment of venous insufficiency and ischemia, for promoting the healing of tendon and ligament injuries, for alleviation of the symptoms of urinary and faecal incontinence.

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 06 May 2016 until 15 April 2021 and remains valid subject to satisfactory surveillance audits.

Re certification audit due before 10 April 2019

Issue 8. Certified since 26 October 2010

Certification is based on reports numbered GB/PC 230088

Authorised by

SGS United Kingdom Ltd, Notified Body 0120

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