

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 16 December 2019 until 15 April 2024
and remains valid subject to satisfactory surveillance audits.

Issue 1. Certified since 26 October 2010

and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered GB/PC 230088

Authorised by



SGS Belgium NV, Notified Body 1639

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LPMD5007 - Certificate CE1639 Annex II-4_EN rev. 02

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