EC Certificate Production Quality Assurance System: Certificate GB19/964574



The management system of

Outside In (Cambridge) Limited also trading as Lumie

3 The Links, Trafalgar Way, Bar Hill, Cambridge, CB23 8UD, UK

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices. Annex V

For the following products

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

This certificate is valid from 16 December 2019until 17 October 2022 and remains valid subject to satisfactory surveillance audits.

Issue 1. Certified since 29 November 1999 and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered GB/PC/ 228934

Authorised by

Pieter Weterings

SGS Belgium NV, Notified Body 1639

SGS House Noorderlaan 87 2030 Antwerp Belgium t+32 (0)3 545-48-48 f+32 (0)3 545-48-49 www.sgs.com

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Outside In (Cambridge) Limited also trading as Lumie

Directive 93/42/EEC

on medical devices, Annex V

Issue 1

Detailed scope

Light therapy devices for the treatment of Seasonal affective disorder (SAD):

LUMIE Arabica LBA

Lumie Desklamp LDL2

LUMIE Zest

• LUMIE Brazil

Lumie Vitamin L LBV

· Lumie Vitamin L PRO LBVP

· Lumie Soleil LBV-S

Light therapy device for the treatment of Acne Vulgaris
• LUMIE CLEAR.

Where the above scope includes class IIb or class III medical device(s), a valid EC Type Examination Certificate according to Annex III is a mandatory requirement for each device in addition to this certificate to place that device on the market