

The management system of

Brymill Cryogenic Systems

105 Windermere Avenue, Ellington, CT, 06029, United States

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

**Cryosurgery Guns for the delivery of Liquid Nitrogen
in the treatment of Cutaneous Skin Lesions.**

**Cryosurgery Closed Probes, Cryochambers, Straight Sprays, Bent Sprays,
Straight Spray Extensions, Bent Spray Extensions Apertures - Accessories
to the Cryosurgery Guns to control the output of the Liquid Nitrogen.
Cryochambers – accessories to the cryosurgery guns to allow surgical
destruction of tissue up to 3 cm deep by causing LN2
to pool continuously over deep lesions.**

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 24 May 2024
and remains valid subject to satisfactory surveillance audits.
Issue 1. Certified since 30 November 1996
and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered WW/MC 06139

Authorised by

Pieter Weterings
Certification Manager

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

LPMD5007 - Certificate CE 1639 Annex II-4, EN rev 02

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