

EC Certificate Full Quality Assurance System: Certificate US19/819943492

The management system of

DGH Technology, Inc.

110 Summit Drive
Suite B, Exton, PA, 19341, 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

Ultrasonic diagnostic equipment used by health care professionals to measure and visualize structures in the human eye, aiding them in the detection, assessment and treatment of conditions affecting vision and patient health.

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 11 August 2023 and remains valid subject to satisfactory surveillance audits.
Issue 1. Certified since 10 August 2006 and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered WW/MC 214712

Authorised by

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LPND5007 - Certificate CE1639 Annex II-4_EN rev. 02

Page 1 of 1

