EC Certificate Full Quality Assurance System: Certificate US06/69508

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 9 October 2015 until 11 August 2020
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 14 July 2018
Issue 7. Certified since 10 August 2006
Certification is based on reports numbered WW/ME 214712

Authorised by

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