

The management system of

Keeler Ltd

Clewer Hill Road, Windsor, Berkshire, SL4 4AA, UK

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex V

For the following products

**Applanation Tonometer to aid diagnosis
and measurement of intraocular pressure**

**Disposable Applanation Tonometer Cone
for use in testing of intraocular pressure**

Disposable Cryo Probe for use with Ophthalmic Surgery Devices

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.

This certificate is valid from 18 May 2021 until 23 March 2023
and remains valid subject to satisfactory surveillance audits.
Issue 2. Certified since 08 September 2009

Certification is based on reports numbered GB/PC/ 240569

Authorised by



Global Medical Devices Head of Notified Body

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

LPMD5008 - Certificate CE1639 AnnexV_EN Rev. 02

Page 1 of 1

