

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**

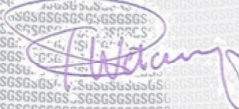
Where the above scope includes class IIb or class III medical device(s), a valid EC Type Examination Certificate according to Annex III is a mandatory requirement for each device in addition to this certificate to place that device on the market.

This certificate is valid from 28 February 2020 until 30 September 2022  
and remains valid subject to satisfactory surveillance audits.

Issue 1. Certified since 08 September 2009  
and first certified by SGS Belgium NV since 28 February 2020

Certification is based on reports numbered GB/PC/ 240569

Authorised by



**SGS Belgium NV, Notified Body 1639**

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LPMD5008 - Certificate CE1639 Annex V, EN rev. 01

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