EC Certificate Production Full Quality Assurance System: Certificate GB20/965237

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