EC Certificate Production 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 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

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