

EC Certificate Production Quality Assurance System: Certificate CN19/41105 The management system of

Ningbo Ming Sing Optical R & D Co., Ltd.

No.702, North Tiantong Road, Yinzhou District, Ningbo, Zhejiang Province, 315192, P.R. China

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex V Restricted to the aspects of manufacture concerned with the conformity of the devices with metrological requirements

For the following products

AUTO REF/KERATOMETER

(Model: KR-9000, KR-9200, KR-9, KR-9300, RM-9000, RM-9200, AR-9, RM-9300, KR-9600, VX90, KR-9800, VX95, RM-9600, RM-9800),

COMPUTERIZED VISION TESTER

(Model: CV-7000, CV-7200, CV-7600, CV-7800, CV-7900, CV55, VX55, CV60, VX60, CV70, VX70)

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 16 December 2019 until 12 November 2023 and remains valid subject to satisfactory surveillance audits. Issue 1. Certified since 05 April 2015 and first certified by SGS Belgium NV since 16 December 2019

> > Certification is based on reports numbered CN/NGB/ 5756 Authorised by

SGS Belgium NV, Notified Body 1639

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