

The management system of

Network Medical Products Ltd

Coronet House, Kearsley Road, Ripon, North Yorkshire, HG4 2SG, UK

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC on medical devices, Annex V

For the following products

**Sterile single use Precision LASIK blades
Sterile single use miniature ENT blades (Netblades)
Sterile single use Coronet Artificial Anterior Chamber
to support donor cornea material prior to implantation.**

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 11 July 2018 until 30 May 2023
and remains valid subject to satisfactory surveillance audits.

Re certification audit due before 30 May 2021

Issue 12. Certified since 27 July 2012

Certification is based on reports numbered GB/PC 230480

Authorised by

SGS United Kingdom Ltd, Notified Body 0120

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