

EC Certificate Full Quality Assurance System: Certificate GB13/88234

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 II (excluding Section 4)

For the following products

- Sterile and non-sterile single use Corneal trephines**
- Sterile single use DCR bodkins**
- Sterile single use canaliculus intubation sets**
- Sterile single use Ventilation tubes and grommets**
- Sterile single use Otological implants for ossicular replacement**
- Sterile single use CORONET corneal tissue delivery system**
- Sterile single use Donor trephine punch**
- Sterile and non-sterile single use aspirating suction tubes and fine ends for the removal of particulates during ENT surgery**
- Sterile single use Epistaxis catheter**

Where the above scope includes class III medical device(s), a valid EC Design Examination Certificate according to Annex II (Section 4) 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 11. Certified since 17 September 1997

Certification is based on reports numbered GB/PC 230480

Authorised by

SGS United Kingdom Ltd, Notified Body 0120

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