EC Certificate Full Quality Assurance System: Certificate GB19/964588

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 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 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

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 16 December 2019 until 30 May 2023
and remains valid subject to satisfactory surveillance audits.
Issue 1. Certified since 17 September 1997
and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered GB/PC 230480

Authorised by

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