

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

Restricted to the aspects of manufacture concerned with securing and maintaining sterile conditions

For the following products

Sterile single use PVA and cellulose sponge products for Ophthalmic and ENT fluid management.

Sterile single use Nasal Splints for airway management support and separation.

Sterile single use Epistaxis catheter.

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 30 May 2023 and remains valid subject to satisfactory surveillance audits.

Issue 1. Certified since 29 June 2006

and first certified by SGS Belgium NV since 16 December 2019

Certification is based on reports numbered GB/PC/ 230480

Authorised by

SGS Belgium NV, Notified Body 1639

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