

EC Certificate Full Quality Assurance System: Certificate US07/740

The management system of

Southmedic Inc

50 Alliance Blvd,
Barrie, Ontario, L4M 5K3, Canada

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

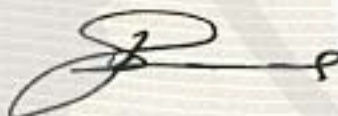
Open oxygen delivery device with carbon dioxide monitoring; Oxygen mask with carbon dioxide monitoring; PVC tubing and silicone tubing, connectors, Nasal cannula, Adult and paediatric face masks (aerosol and oxygen); Tracheostomy oxygen masks; Sterile and non-sterile surgical blades, sterile and non-sterile disposable and safety scalpels.

For placing on the market of Class III devices covered by this certificate, an EC Design Examination Certificate according to Annex II (Section 4) is required.

This certificate is valid from 2 December 2011 until 9 January 2015
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 8 December 2012
Issue 8. Certified since 10 January 2007

Certification is based on reports numbered WW/PCI 215244

Authorised by



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