

EC Certificate Production Quality Assurance System: Certificate
MY12/00920

The management system of

Mediplas Respiratory Products Sdn. Bhd.

No. 7, Jalan KIP 3
Taman Perindustrian KIP, 52200 Kuala Lumpur
MALAYSIA

has been assessed and certified as meeting the requirements of

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

For the following products

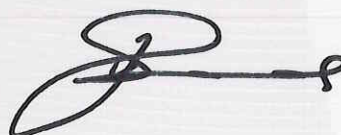
**Breathing circuits (including sterile: Tubing extension Sets, Suction
Circuit Tubing, Oxygen Tubing), Fittings and Adaptors and Sterile
Laryngoscopes.**

For placing on the market of Class IIb or Class III devices covered by this certificate, an EC Type
Examination Certificate according to Annex III is required.

This certificate is valid from 30 June 2012 until 30 June 2015 and remains
valid subject to satisfactory surveillance audits.
Re certification audit due before 30 April 2015
Issue 2. Certified since 02 May 2012

Certification is based on reports numbered MY/KUL MY02884

Authorised by



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