EC Certificate Full Quality Assurance System: Certificate US98/15042

The management system of

CPR Medical Devices Inc.

161 Don Park Road,
Markham, Ontario, L3R IC2, 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

OXYLATOR® EM-100, FR-300, EMX and HD resuscitation equipment.

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 27 April 2016 until 10 June 2020
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 10 February 2018
Issue 10. Certified since 23 November 1998

Certification is based on reports numbered WW/MC 09564

Authorised by

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