EC Certificate Production Quality Assurance System: Certificate GB01/53125

The management system of

Medfor Products Ltd

Unit 2, Gresham Industrial Estate, Eastern Road,
Aldershot, Hampshire, GU12 4YD, UK

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex V

For the following products

Sterile and non sterile containers for the transit and storage of human organs and tissues prior to transplantation.

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 18 February 2016 until 19 December 2020
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 19 December 2018
Issue 8. Certified since 17 September 2001

Certification is based on reports numbered GB/PC 200659

Authorised by

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