EC Certificate Production Quality Assurance System: Certificate GB14/91301

The management system of

Al-Shifa Medical Syringe Manufacturing Co.

1st Industrial City, PO Box 7917, Dammam,
Kingdom of Saudi Arabia

has been assessed and certified as meeting the requirements of

Directive 93/42/EEC

on medical devices, Annex V

For the following products

- Sterile, Single-Use Hypodermic Syringes with needles.
- Sterile, Single-Use U-100 Insulin Syringes, Fixed needle, with and without protective end cap.

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 07 May 2014 until 24 April 2019 and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 01 December 2016
Issue 2. Certified since 24 April 2014

Certification is based on reports numbered GB/PI 232016

Authorised by

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