EC Certificate Full Quality Assurance System: Certificate GB15/92506

The management system of

Mediplus (India) Limited

1261-1262, M.I.E. Part B, Bahadurgarh-124507, (Haryana), India

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

Sterile I. V. Cannula, Sterile I.V. Cannula with Safety Feature, Sterile 3-Way Stop Cock, Sterile 3-Way Stop Cock with Extension Tubing, Sterile Injection Stopper, Sterile Male- Female Luer Lock, Sterile Luer Lock, Sterile A.V. Fistula Needle / Set & Sterile Pressure Monitoring Line.

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 07 April 2015 until 12 January 2020 and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 16 September 2017
Issue 2. Certified since 12 January 2015

Certification is based on reports numbered GB/PI 233293

Authorised by

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