EC Certificate Full Quality Assurance System: Certificate IN15/92506

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.

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 01 April 2019 until 12 January 2023
and remains valid subject to satisfactory surveillance audits.
Re certification audit due before 16 September 2020
Issue 2. Certified since 12 January 2015

Certification is based on reports numbered IN/GUR 235607

Authorised by

Jonathan M. Hall

SGS United Kingdom Ltd, Notified Body 0120
202B Worle Parkway, Weston-super-Mare, BS22 6WA UK
t +44 (0)1934 522917 f +44 (0)1934 522137 www.sgs.com

SGS CE 02 0215