EC Certificate Full Quality Assurance System: Certificate IN19/818843707

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 Needle Free Valve and Sterile Pressure Monitoring line/ Sterile Extension Tube

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 09 April 2020 until 12 January 2023
and remains valid subject to satisfactory surveillance audits.
Issue 2. Certified since 12 January 2015
and first certified by SGS Belgium NV since 16 December 2019.

Certification is based on reports numbered IN/GUR 235607

Authorised by

Digitally signed by Talpiş Andrei
Date: 2021.09.14 20:15:54 EEST
Reason: MoldSign Signature
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