CERTIFICATE OF EUROPEAN UNION AUTHORISED REPRESENTATIVE

CE

This is to certify that Translite LLC.

has duly registered the following relevant product types with the UK Competent Authority through its Appointed Representative in accordance with *Article 14* of the Council Directive 93/42/EEC as revised by Council Directive 20/47/EC concerning medical devices (The "Medical Devices Directive") (UK Medical Devices Regulation 1994: Regulation 14).

********** Applicable ANNEX **********

Annex VII

********** Scope of Supply **********

Product Class I non-sterile. Product Family: Veinlite

Competent Authority Registration Reference CA008352

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In accordance by self-declaration with *Article 11* and *Annex VII* for Class I devices may apply the CE Mark

********** Appointment **********

We certify that M Devices Group/EC Rep Ltd was appointed as the Authorised Representative on the 4th August 2004

Signature Authorised Representative	Date: 7 Sept 2018

Certificate No. MDG—AR-40 Valid to 4-August-2019

EC REP Ltd

Health & Education Centre, The Church
Portland Street, Southport PR8 1HU
United Kingdom