CERTIFICATE OF EUROPEAN UNION AUTHORISED REPRESENTATIVE

This is to certify that TransLite LLC *US-MF-000029360*

CE

345 Commerce Green Blvd, Sugar Land, TX 77478, USA

has duly registered as a manufacturer as a result of Brexit, with the Irish Competent Authority through its Appointed Authorised Representative in accordance with *Article 14* of the Council Directive 93/42/EEC concerning medical devices (The "Medical Devices Directive") and Medical Devices Regulation 2017/745.

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

Annex VII E C Declaration of Conformity,

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

BSI QMS ISO 13485:2016

Product Class I non-sterile. Product Family: Veinlite

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Registered reference with the HPRA: E C Rep Limited / Translite LLC - MDD - IE/CA01/R/GM/1404

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

We certify that E C Rep Ltd was appointed as the Authorised Representative on the 6 May 2019

Signature
Authorised Representative

Date: 26 Feb 2024

E C Rep Ltd, 5 Fitzwilliam Square East, Dublin, D02 R744, Ireland

Certificate No. AR-T-41 **Valid to 5 May 2025**