

CERTIFICATE OF SWITZERLAND AUTHORISED REPRESENTATIVE



This is to certify that TransLite LLC.

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

Under the regulations MedDO with the Swiss Competent Authority through its Appointed Authorised Representative in accordance with (Art. 4 para. 1 let. g MedDO, Art. 4 para. 1 let. f IvDO and Article 11 REGULATION (EU) 2017/745 OF THE EUROPEAN PARLIAMENT AND OF THE COUNCIL of 5 April 2017 (Medical Devices Regulation 2017/745) MDR.

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

Annex VII Declaration of Conformity,

SRN US-MF-000029360

BSI QMS ISO 13485:2016 Exp Date: 2024-09-01 No. FM 633388

Product Class I non-sterile. Product Family: Veinlite

Registered under the transition regulations

We certify that E C Rep GmbH (UID CHE-353.390.735) was appointed as the Authorised Representative on the 15 March 2023

Signature	
Authorised	Representative



Date: 15 Mar 2023



E C Rep GmbH, Bahnhofstrasse 71, 8001, & Luegislandstrasse 105, 8051 Zurich

Certificate No. CH-AR-0 Valid to 30 June 2024