PRODUCT IDENTIFICATION	
Device Family	**Device trade names**
LightForce Therapy Lasers	LIghtForce FXi LightForce EXPi LightForce XPi LightForce XLi

MANUFACTURER		
Name of company	**Address**	**Representative**
LiteCure, LLC	101 Lukens Dr, Suite A New Castle, DE 19720 USA	Quality & Regulatory Director

AUTHORIZED REPRESENTATIVE		
Name of company	**Address**	**Contact**
Emergo Europe	Prinsessegracht 20 2514 AP The Hague The Netherlands	P: +31 (0)70 345 85 70 F: +31 (0) 70 346 72 99

REGISTRATION INFORMATION		
Notified Body and ID #	**Marking**	**CE certificate number**
BSI (NL) – 2797	CE 2797	CE 542523

CONFORMITY ASSESSMENT		
Device classification	**Route to compliance**	**Standards applied**
Class IIa Rule 9	Annex II of MDD 93/42/EEC Council Directive	IEC 60601-1:2005+A1:2012 IEC 60601-1-2:2015 IEC 60601-1-6:2010+A1:2013 IEC 60601-2-22:2007+A1:2013 IEC 60825-1:2014

Litecure, LLC declares that the above-mentioned products meet the provision of the Council Directive 93/42/EEC for Medical Devices and amendment 2007/47/EC as transposed in the national laws of the Member States.

Eric Rock	
Quality and Regulatory Representative *Print Name*	*Sign*
New Castle, Delaware, USA	**25 MAY 2021**
Place	*Date*