3Gen Inc

31521 Rancho Viejo Rd, Suite #104
San Juan Capistrano, CA 92675, USA

Declaration of Conformity

Annex of the Medical Device Directive under which Declaration is made: **VII**

Date of Issuance: September 16, 2018 (Supersedes Declaration dated: May 1, 2018)

Manufacturer:
3Gen, Inc.
31521 Rancho Viejo Rd., #104,
San Juan Capistrano, CA 92675

EC Representative:
M. Devices Group
Healthcare Education Centre,
The Church, Portland Street, Southport, PR8 1HU, UK

This Declaration is applicable to all products listed and manufactured after the Date of Issuance of this Declaration of Conformity.

We hereby declare under our sole responsibility, that the following products, comply fully with the requirements of the Medical Device Directive (93/42/EEC, amended by directives 98/79/EC, 2000/70/EC and 2007/47/EC).

Model Number	Device Class
DermLite DL100	I
DermLite Platinum	I
DermLite Pro DP-R	I
DermLite Foto	I
DermLite II Pro (DL2Pro)	I
DermLite II Multispectral (DL2MS)	I
SkinLite ALT100-0033	I
DermLite II Pro HR (DL2HR)	I
DermLite II Fluid	I
DermLite II Hybrid (DL2HM)	I
DermLite Lumio (LUM)	I
SkinLite II	I
DermLite Carbon (DermLite DLC)	I
Alumina	I
DermLite 3 (DL3)	I
DermLite 1 (DL1)	I
DermLite Lumio S (DLUS)	I
DermLite Cam (DLCAM)	I
DermLite 3 (DL3N)	I
DermLite Foto II Pro (Foto II Pro)	I
DermLite Lumio UV (LUM UV)	I
DermLite 1 Basic (DL1B)	I
DermLite DL4 (DL4)	I
DermLite HUD (HUD-I6)	I

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Model Number (Continued)	Device Class (Continued)
DermLite DL200 Hybrid	I
DermLite DL200 HR	I
DermLite DL4W	I
DermLite GL	I
DermLite DL200 Hybrid Gold	I
DermLite Foto X	I

Signed:

John Bottjer, President, 3Gen Inc.