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: March 10, 2015 (Supersedes Declaration dated: September 10, 2014)

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	I
DermLite II Fluid	I
DermLite II Hybrid	I
Lumio	I
SkinLite II	I
Carbon	I
Alumina	I
DermLite 3 (DL3)	I
DermLite 1 (DL1)	I
Lumio S	I
DLCam	I
DL3N	I
Foto II Pro (DLF2-PRO)	I
Lumio UV	I
DermLite 1 Basic (DL1B)	I
DermLite 4 (DL4)	I

Signed:

John Bottjer, President, 3Gen Inc.

Document # 000-0006, Rev 17.