

## Declaration of Conformity

to Annex VII of the EC Medical Device Directive

Date: March 2, 2009

We **3Gen, LLC**, located at 31521 Rancho Viejo Rd., #104, San Juan Capistrano, CA 92675, USA 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 and 2000/70/EC).

**Product:** Epiluminescence device

<b>Model Number</b>	<b>Starting Serial Number</b>
DermLite DL100	DLA01200300000
DermLite Platinum	DLP01200300000
DermLite Pro DP-R	DPR02200300000
DermLite Foto	DLF02200300000
DermLite II Pro (DL2Pro)	DL2PA200400000
DermLite II Multispectral (DL2MS)	DL2MA200400000
SkinLite ALT100-0033	SKN01200400001
DermLite II Pro HR	D2H02200500001
DermLite II Fluid	DLF02200600001
DermLite II Hybrid	D2A02200600001
Lumio	LUM02200700001
SkinLite II	SKN02200700001
Carbon	DLC01200800001
Alumina	DAL02200800001
DermLite 3	DL301200900001

**Class:** Class I (according to Annex IX of the directive)

Signed:



John Bottjer, President, 3Gen LLC