CERTIFICATE OF EUROPEAN UNION AUTHORISED REPRESENTATIVE

CE

This is to certify that 3 Gen, Inc.

has duly registered the following relevant product types with the UK Competent Authority through its Appointed Representative in accordance with *Article 14* of the Council Directive 93/42/EEC (revised by 2007/47/EC) concerning medical devices

(The "Medical Devices Directive") (UK Medical Devices Regulation 1994: Regulation 14).

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

Annex VII

********** Scope of Supply **********

DERMLITE Class I non-sterile

****

In accordance by self-declaration with *Article 11* and *Annex VII* for Class I devices may apply the CE Mark

********** Appointment **********

We certify that E C Rep Ltd/ M. Devices Group was appointed as the Authorised Representative on the 15th Nov 2003

Signature
Authorised Representative

Date:
15 November 2018

Certificate No. MDG-1034-AR Valid to 15 Nov 2019

Healthcare Education Centre,
Portland Street, Southport, PR8
1HU, England