CERTIFICATE OF EUROPEAN UNION AUTHORISED REPRESENTATIVE

CE

This is to certify that 3 Gen LLC.

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

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

Annex VII

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

DemLite - Class I non-sterile

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

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

We certify that M. Devices Group was appointed as the Authorised Representative on the 15th Nov 2003

MHRA Registration Reference CA 008007 11-Dec-2003

Signature
Authorised Representative

Paul Bennett

Date 04Nov2009

Certificate No. MDG-1034-AR **Valid to 14 Nov 2012**

MDEVICES group

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