Quality System Approval Certificate

Medical Devices Directive 93/42/EEC

*The National Standards Authority of Ireland as a duly designated Notified Body, (identification number **0050**), for the purposes of the European Communities (Medical Devices) Regulations (S.I. No. 252 of 1994)*

APPROVES THE QUALITY SYSTEM APPLIED BY

Sunlight Medical, Inc.

5570 Florida Mining BLVD S.
Suite 603
Jacksonville
FL 32257
USA

to the Product Family

Mechanical Pipette (In Vitro Fertilization Pipette)

GMDN Code: 38522

on the basis of examination under the requirements of Directive 93/42/EEC on Medical Devices Annex V.
*The use of the NSAI Notified Body identification number **0050** in conjunction with CE Marking of Conformance for this product family is hereby authorized.*

Registration Number:	**252.944**
Original Approval:	**24 September 2014**
Last Amended on:	**31 May 2017**
Remains valid until:	**23 September 2020**

Signed:

Approved by:
Geraldine Larkin
Chief Executive Officer, NSAI

Approved by:
Susan Murphy
European Medical Device Operations Manager

This certificate remains valid on condition that the Approved Quality System is maintained in an adequate and efficacious manner .
Details of the current product range and operational locations included within the scope of this approval can be obtained from NSAI

National Standards Authority of Ireland, 1 Swift Square, Northwood, Santry, Dublin 9, Ireland.