



NSAI

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:	30 April 2019
Remains valid until:	23 September 2022

Signed:

Approved by:
Geraldine Larkin
Chief Executive Officer, NSAI

Approved by:
Dr Caroline Dore Geraghty
Director, Medical Devices

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.

