

**K072600 INTRACYTOPLASMIC SPERM AND SPERMATID
INJECTION, HOLDING, ZONE DRILLING, PARTIAL ZONE
DISSECTION AND DENUDING PIPETTES**May 19, 2008
248 days to decisionK072600 · Product code: **MQH** · Obstetrics & Gynecology
Source: <https://www.510kdatabase.net/k072600/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Microtools, Assisted Reproduction (pipettes) (MQH)
Date received	Sep 14, 2007
Decision date	May 19, 2008
Days to decision	248 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Sunlight Medical, Inc.
Location	Jacksonville, FL, US
Contact	DUNSONG YANG
510(k) history	2 submissions · 2 cleared · 2008-2010

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k072600/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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