

**K022643 BLASTOMERE BIOPSY PIPETTE, MODELS H-55530
AND H-55540**Nov 5, 2002
89 days to decisionK022643 · Product code: **MQH** · Obstetrics & Gynecology
Source: <https://www.510kdatabase.net/k022643/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Microtools, Assisted Reproduction (pipettes) (MQH)
Date received	Aug 8, 2002
Decision date	Nov 5, 2002
Days to decision	89 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Swemed Lab International AB
Location	Washington, DC, US
Contact	HOWARD M HOLSTEIN
510(k) history	4 submissions · 4 cleared · 1999-2006

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k022643/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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