

**K122337 OVUSENSE**Aug 6, 2013  
369 days to decisionK122337 · Product code: **LHD** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k122337/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Fertility Diagnostic, Proceptive (LHD)
Date received	Aug 2, 2012
Decision date	Aug 6, 2013
Days to decision	369 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Fertility Focus, Ltd.</b>
Location	Swanee, GA, US
Contact	PENNY NORTHCUTT
510(k) history	1 submissions · 1 cleared · 2013-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k122337/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated June 8, 2026