

K050094 LADY COMP USAApr 20, 2006
461 days to decisionK050094 · Product code: **LHD** · Obstetrics & Gynecology
Source: <https://www.510kdatabase.net/k050094/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Fertility Diagnostic, Proceptive (LHD)
Date received	Jan 14, 2005
Decision date	Apr 20, 2006
Days to decision	461 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Lady Comp USA
Location	Jeffersonville, IN, US
Contact	MICHAEL CARTAIN
510(k) history	1 submissions · 1 cleared · 2006-2006

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