FDA 510(k) Premarket Notification

K050094 LADY COMP USA

CLEARED

Apr 20, 2006
461 days to decision

K050094 · 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