

K870999 GENDER CHOICE INFORMATION KIT, FEMALE AND MALEJul 14, 1987
125 days to decisionK870999 · Product code: **LHD** · Obstetrics & Gynecology
Source: <https://www.510kdatabase.net/k870999/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Fertility Diagnostic, Proceptive (LHD)
Date received	Mar 11, 1987
Decision date	Jul 14, 1987
Days to decision	125 days
Third-party review	No

APPLICANT

Company	Procare C/O Wiley, Rein & Fielding
Location	Washington, DC, US
Contact	ANDREW S KRULWICH
510(k) history	1 submissions · 1 cleared · 1987-1987

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