FDA 510(k) Premarket Notification

K870999 GENDER CHOICE INFORMATION KIT, FEMALE AND MALE

CLEARED

Jul 14, 1987
125 days to decision

K870999 · 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