FDA 510(k) Premarket Notification

K810832 FORECASTER

CLEARED

Jun 24, 1981
91 days to decision

K810832 · Product code: **LHD** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k810832/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Fertility Diagnostic, Proceptive (LHD)
Date received	Mar 25, 1981
Decision date	Jun 24, 1981
Days to decision	91 days
Third-party review	No

APPLICANT

Company	**Forecaster Co.**
Location	Mchenry, IL, US
510(k) history	1 submissions · 1 cleared · 1981-1981