FDA 510(k) Premarket Notification

K860233 5 MHZ INTRACAVITARY PROBE, MODEL H4222V

CLEARED

Jun 2, 1986
131 days to decision

K860233 · Product code: **HEM** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k860233/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Imager, Ultrasonic Obstetric-gynecologic (HEM)
Date received	Jan 22, 1986
Decision date	Jun 2, 1986
Days to decision	131 days
Third-party review	No

APPLICANT

Company	**General Electric Co.**
Location	Mchenry, IL, US
Contact	HOWARD II
510(k) history	254 submissions · 254 cleared · 1976-2011