FDA 510(k) Premarket Notification

K831212 ENDOSCANN

CLEARED

Jun 15, 1984
428 days to decision

K831212 · Product code: **HFF** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k831212/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Aspirator, Endometrial (HFF)
Date received	Apr 14, 1983
Decision date	Jun 15, 1984
Days to decision	428 days
Third-party review	No

APPLICANT

Company	**Axcan Scientific Corp.**
Location	Walker, MI, US
510(k) history	5 submissions · 5 cleared · 1982-1984