FDA 510(k) Premarket Notification

K873041 A-V RECORDER

CLEARED

Feb 4, 1988
184 days to decision

K873041 · Product code: **DPT** · Cardiovascular
Source: https://www.510kdatabase.net/k873041/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Probe, Blood-flow, Extravascular (DPT)
Date received	Aug 4, 1987
Decision date	Feb 4, 1988
Days to decision	184 days
Third-party review	No

APPLICANT

Company	**Life Sciences Manufacturing, Inc.**
Location	Mchenry, IL, US
Contact	JAMES H VEZINA
510(k) history	12 submissions · 12 cleared · 1979-1991