FDA 510(k) Premarket Notification

K955802 PATHFINDER

CLEARED

Jan 29, 1997
404 days to decision

K955802 · Product code: **DRF** · Cardiovascular
Source: https://www.510kdatabase.net/k955802/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Dec 22, 1995
Decision date	Jan 29, 1997
Days to decision	404 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Cardima, Inc.**
Location	Fremont, CA, US
Contact	GABRIEL VEGH
510(k) history	12 submissions · 12 cleared · 1993-2006