FDA 510(k) Premarket Notification

CLEARED

K971975 CARDIMA PATHFINDER 1.5F MAPPING MICROCATHETER

Jul 1, 1998
398 days to decision

K971975 · Product code: **DRF** · Cardiovascular
Source: https://www.510kdatabase.net/k971975/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	May 29, 1997
Decision date	Jul 1, 1998
Days to decision	398 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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