

**K971975 CARDIMA PATHFINDER 1.5F MAPPING
MICROCATHETER**Jul 1, 1998
398 days to decisionK971975 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k971975/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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