

**K983653 RADII DIAGNOSTIC ELECTROPHYSIOLOGY
CATHETER, MODEL # 30411 B, C AND D**Dec 22, 1998
64 days to decisionK983653 · Product code: DRF · Cardiovascular
Source: <https://www.510kdatabase.net/k983653/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Oct 19, 1998
Decision date	Dec 22, 1998
Days to decision	64 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Cardiac Pathways Corp.
Location	Sunnyvale, CA, US
Contact	ERIN DIGNAN
510(k) history	5 submissions · 4 cleared · 1997-2000

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k983653/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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