

**K992912 CARDIAC PATHWAYS REFERENCE CATHETER AND  
REFERENCE CATHETER WITH TRACKING, CARDIAC  
PATHWAYS RADII CATHETER AND RADII CATH**Mar 8, 2000  
191 days to decisionK992912 · Product code: **DRF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k992912/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Aug 30, 1999
Decision date	Mar 8, 2000
Days to decision	191 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Cardiac Pathways Corp.</b>
Location	Sunnyvale, CA, US
Contact	DEBRA S ECHT
510(k) history	5 submissions · 4 cleared · 1997-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k992912/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated July 4, 2026