FDA 510(k) Premarket Notification

CLEARED

K983653 RADII DIAGNOSTIC ELECTROPHYSIOLOGY CATHETER, MODEL # 30411 B, C AND D

Dec 22, 1998
64 days to decision

K983653 · 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