

K940168 DIAGNOSTIC II CATHETERJul 27, 1994
201 days to decisionK940168 · Product code: **DRF** · CardiovascularSource: <https://www.510kdatabase.net/k940168/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Jan 7, 1994
Decision date	Jul 27, 1994
Days to decision	201 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Ep Technologies, Inc.
Location	Mountain View, CA, US
Contact	ALAN MARQUARDT
510(k) history	15 submissions · 15 cleared · 1988-2005

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k940168/> Data sourced from FDA 510(k) public records (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. - Generated July 2, 2026