FDA 510(k) Premarket Notification

K940168 DIAGNOSTIC II CATHETER

CLEARED

Jul 27, 1994
201 days to decision

K940168 · Product code: **DRF** · Cardiovascular
Source: 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