FDA 510(k) Premarket Notification

K924109 STEEROCATH II

CLEARED

Dec 16, 1993
489 days to decision

K924109 · Product code: **DRF** · Cardiovascular
Source: https://www.510kdatabase.net/k924109/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Aug 14, 1992
Decision date	Dec 16, 1993
Days to decision	489 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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