

**K924109 STEEROCATH II**Dec 16, 1993  
489 days to decisionK924109 · Product code: **DRF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k924109/>**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 14, 1992
Decision date	Dec 16, 1993
Days to decision	489 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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

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