

K900765 STEEROCATH(TM)May 7, 1990
80 days to decisionK900765 · Product code: **DRF** · Cardiovascular
Source: <https://www.510kdatabase.net/k900765/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Feb 16, 1990
Decision date	May 7, 1990
Days to decision	80 days
Third-party review	No

APPLICANT

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

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