

**K900452 QUATTRO CATHETER**Mar 14, 1990  
43 days to decisionK900452 · Product code: **DRF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k900452/>**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	Jan 30, 1990
Decision date	Mar 14, 1990
Days to decision	43 days
Third-party review	No

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k900452/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated July 3, 2026