

**K984150 PROCATH ELECTROPHYSIOLOGY CATHETER, EP  
MEDSYSTEMS ELECTROPHYSIOLOGY CATHETER**

Feb 22, 1999  
95 days to decision

K984150 · Product code: DRF · Cardiovascular  
Source: <https://www.510kdatabase.net/k984150/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	Nov 19, 1998
Decision date	Feb 22, 1999
Days to decision	95 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Ep Medsystems</b>
Location	West Berlin, NJ, US
Contact	JOSEPH C GRIFFIN, III
510(k) history	12 submissions · 12 cleared · 1996-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k984150/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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