

**K894500 DAIG ELECTROPHYSIOLOGY CATHETER**Sep 18, 1989  
61 days to decisionK894500 · Product code: **DRF** · CardiovascularSource: <https://www.510kdatabase.net/k894500/>**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	Jul 19, 1989
Decision date	Sep 18, 1989
Days to decision	61 days
Third-party review	No

**APPLICANT**

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Company	<b>Daig Corp.</b>
Location	Mchenry, IL, US
Contact	J FLEISCHHACKER
510(k) history	63 submissions · 63 cleared · 1977-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k894500/>, 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