

**K942379 DAIG DIAGNOSTIC ELECTROPHYSIOLOGY
CATHETER**Dec 27, 1994
232 days to decisionK942379 · Product code: **DRF** · Cardiovascular
Source: <https://www.510kdatabase.net/k942379/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Catheter, Electrode Recording, Or Probe, Electrode Recording (DRF)
Date received	May 9, 1994
Decision date	Dec 27, 1994
Days to decision	232 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k942379/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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