

**K875059 DISPOSABLE BIPOLAR TEMPORARY PACING  
ELECTRODE**Feb 12, 1988  
65 days to decisionK875059 · Product code: **LDF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k875059/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Dec 9, 1987
Decision date	Feb 12, 1988
Days to decision	65 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/k875059/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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