

**K800352 PACEMAKER, ELECTRODE**May 28, 1980  
103 days to decisionK800352 · Product code: **DTB** · CardiovascularSource: <https://www.510kdatabase.net/k800352/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Feb 15, 1980
Decision date	May 28, 1980
Days to decision	103 days
Third-party review	No

**APPLICANT**

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Company	<b>Daig Corp.</b>
Location	Mchenry, IL, US
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/k800352/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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