

**K802552 PERM. ENDOCARDIAL UNIPOLAR TINED/WEDGE**Jan 16, 1981  
93 days to decisionK802552 · Product code: **DTB** · CardiovascularSource: <https://www.510kdatabase.net/k802552/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Oct 15, 1980
Decision date	Jan 16, 1981
Days to decision	93 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/k802552/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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