

**K874183 ACTIFIX (R) BIPOLAR ENDOCARDIAL SCREW-IN LEAD**Jan 29, 1988  
108 days to decisionK874183 · Product code: **DTB** · Cardiovascular  
Source: <https://www.510kdatabase.net/k874183/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Oct 13, 1987
Decision date	Jan 29, 1988
Days to decision	108 days
Third-party review	No

**APPLICANT**

---

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

---

**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k874183/> Data sourced from FDA 510(k) public records (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. - Generated July 3, 2026