

**K885240 MODIFIED ANCOR & HI-FLEX UNIPOLAR ACTIVE FIX.
LEAD**Feb 10, 1989
51 days to decisionK885240 · Product code: **DTB** · Cardiovascular
Source: <https://www.510kdatabase.net/k885240/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Dec 21, 1988
Decision date	Feb 10, 1989
Days to decision	51 days
Third-party review	No

APPLICANT

Company	Tpl-Cordis, Inc.
Location	Suffield, CT, US
Contact	DUANE A SCHULTZ
510(k) history	6 submissions · 6 cleared · 1987-1989

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k885240/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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