

**K792604 LIFELINE TLE ENDOCARDIAL PACING LEAD,**Dec 31, 1979  
13 days to decisionK792604 · Product code: **DTB** · CardiovascularSource: <https://www.510kdatabase.net/k792604/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Permanent Pacemaker Electrode (DTB)
Date received	Dec 18, 1979
Decision date	Dec 31, 1979
Days to decision	13 days
Third-party review	No

**APPLICANT**

---

Company	<b>Intermedics, Inc.</b>
Location	Mchenry, IL, US
510(k) history	211 submissions · 201 cleared · 1977-1996

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k792604/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 28, 2026