

**K801800 UNIVERSAL TEMP. PACING ELECT. #080-10000**Oct 23, 1980  
86 days to decisionK801800 · Product code: **LDF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k801800/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Jul 29, 1980
Decision date	Oct 23, 1980
Days to decision	86 days
Third-party review	No

**APPLICANT**

---

Company	<b>Medical Testing System, Inc.</b>
Location	Mchenry, IL, US
510(k) history	5 submissions · 5 cleared · 1977-1981

---

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

Device record: <https://www.510kdatabase.net/k801800/> 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 June 29, 2026