

**K801799 ATRIAL/VENTRICULAR SEQUENTIAL #080-20000**Jan 2, 1981  
157 days to decisionK801799 · Product code: **LDF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k801799/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Jul 29, 1980
Decision date	Jan 2, 1981
Days to decision	157 days
Third-party review	No

**APPLICANT**

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

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

Device record: <https://www.510kdatabase.net/k801799/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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