

**K820003 CARDIOVASCULAR TEMP. PACEMAKER ELEC**Mar 26, 1982  
81 days to decisionK820003 · Product code: **LDF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k820003/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Jan 4, 1982
Decision date	Mar 26, 1982
Days to decision	81 days
Third-party review	No

**APPLICANT**

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Company	<b>Diag Corp.</b>
Location	Walker, MI, US
510(k) history	9 submissions · 9 cleared · 1980-1999

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Device record: <https://www.510kdatabase.net/k820003/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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