

**K855139 FLEXON TEMPORARY CARDIAC CONDUCTIVE  
MULTIFILAMENT**Feb 26, 1986  
65 days to decisionK855139 · Product code: **LDF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k855139/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	Dec 23, 1985
Decision date	Feb 26, 1986
Days to decision	65 days
Third-party review	No

**APPLICANT**

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Company	<b>Davis &amp; Geck, Inc.</b>
Location	Mchenry, IL, US
Contact	PELLERTIER
510(k) history	45 submissions · 42 cleared · 1981-1999

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k855139/>; 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 22, 2026