

**K821518 MYO/CLIP**Jul 8, 1982  
48 days to decisionK821518 · Product code: **LDF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k821518/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Electrode, Pacemaker, Temporary (LDF)
Date received	May 21, 1982
Decision date	Jul 8, 1982
Days to decision	48 days
Third-party review	No

**APPLICANT**

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Company	<b>Alto Development Corp.</b>
Location	Mchenry, IL, US
510(k) history	10 submissions · 10 cleared · 1982-1996

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Device record: <https://www.510kdatabase.net/k821518/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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