

**K982896 ALEXANDER MANUFACTURING CO.  
RECHARGEABLE BATTERY PART NUMBER LP9**Oct 21, 1998  
65 days to decisionK982896 · Product code: **LDD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k982896/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - ST
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Aug 17, 1998
Decision date	Oct 21, 1998
Days to decision	65 days
Third-party review	No

**APPLICANT**

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Company	<b>Alexander Mfg. Co.</b>
Location	Mason City, IA, US
Contact	KEN HEIMENDINGER
510(k) history	39 submissions · 37 cleared · 1997-1998

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k982896/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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