

**K935674 LIFEPAK 9 DEFIBRILLATOR/MONITOR**Dec 16, 1993  
8 days to decisionK935674 · Product code: **LDD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k935674/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dc-defibrillator, Low-energy, (including Paddles) (LDD)
Date received	Dec 8, 1993
Decision date	Dec 16, 1993
Days to decision	8 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Physio-Control Corp.</b>
Location	Mchenry, IL, US
Contact	SHERRI L POCOCK
510(k) history	80 submissions · 78 cleared · 1976-1999

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k935674/> 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 27, 2026