

**K905510 LIFEPAK 9 DEFIBRILLATOR & CARDIAC  
MONITOR/MODIFIED**Dec 20, 1990  
13 days to decisionK905510 · Product code: **LDD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k905510/>**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 7, 1990
Decision date	Dec 20, 1990
Days to decision	13 days
Third-party review	No

**APPLICANT**

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Company	<b>Physio-Control Corp.</b>
Location	Mchenry, IL, US
Contact	D WILLINGHAM
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/k905510/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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