

**K983393 LIFEPAK 500 BIPHASIC**May 5, 1999  
222 days to decisionK983393 · Product code: **MKJ** · CardiovascularSource: <https://www.510kdatabase.net/k983393/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - ST
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Sep 25, 1998
Decision date	May 5, 1999
Days to decision	222 days
Third-party review	No

**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/k983393/>; 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