

**K102972 LIFEPAK 12**Dec 22, 2010  
77 days to decisionK102972 · Product code: **MKJ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k102972/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Oct 6, 2010
Decision date	Dec 22, 2010
Days to decision	77 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Physio-Control, Inc.</b>
Location	Redmond, WA, US
Contact	MICHELLE ACKERMANN
510(k) history	14 submissions · 14 cleared · 1984-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k102972/> 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 26, 2026