

**K012030 PORTABLE INTENSIVE CARE UNIT**Sep 24, 2001  
88 days to decisionK012030 · Product code: **MKJ** · CardiovascularSource: <https://www.510kdatabase.net/k012030/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Jun 28, 2001
Decision date	Sep 24, 2001
Days to decision	88 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Medical Research Laboratories, Inc.</b>
Location	Mchenry, IL, US
Contact	JOEL ORLINSKY
510(k) history	19 submissions · 15 cleared · 1981-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k012030/> 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 25, 2026