

**K041892 ZOLL AED PRO EXTERNAL DEFIBRILLATOR**Feb 4, 2005  
206 days to decisionK041892 · Product code: **MKJ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k041892/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Jul 13, 2004
Decision date	Feb 4, 2005
Days to decision	206 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>ZOLL Medical Corporation</b>
Location	Chelmsford, MA, US
Contact	SEAN REYNOLDS
510(k) history	30 submissions · 30 cleared · 2005-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k041892/> 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 June 5, 2026