

**K072923 ZOLL E SERIES DEFIBRILLATOR**Apr 24, 2008  
192 days to decisionK072923 · Product code: **MKJ** · CardiovascularSource: <https://www.510kdatabase.net/k072923/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Oct 15, 2007
Decision date	Apr 24, 2008
Days to decision	192 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Zoll Medical Corporation, World Wide Headquarters</b>
Location	Chelmsford, MA, US
Contact	SEAN REYNOLDS
510(k) history	21 submissions · 21 cleared · 2007-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k072923/> 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