

**K111594 ZOLL E SERIES**Aug 17, 2011  
71 days to decisionK111594 · Product code: **MKJ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k111594/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Jun 7, 2011
Decision date	Aug 17, 2011
Days to decision	71 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	CHUCK KOLIFRATH
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/k111594/> 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