

**K110168 ZOLL E SERIES WITH 2010 AHA GUIDLINES  
SOFTWARE UPDATE**Feb 17, 2011  
28 days to decisionK110168 · Product code: **MKJ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k110168/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Jan 20, 2011
Decision date	Feb 17, 2011
Days to decision	28 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	EILEEN M BOYLE
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/k110168/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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