

**K110361 ZOLL R SERIES WITH 2010 AHA GUIDELINES
SOFTWARE UPDATE**

Mar 8, 2011
29 days to decision

K110361 · Product code: **MKJ** · Cardiovascular
Source: <https://www.510kdatabase.net/k110361/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Feb 7, 2011
Decision date	Mar 8, 2011
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Zoll Medical Corporation, World Wide Headquarters
Location	Chelmsford, MA, US
Contact	EILEEN M BOYLE
510(k) history	21 submissions · 21 cleared · 2007-2015

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k110361/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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