

**K032439 ZOLL M SERIES WITH RECTILINEAR BI-PHASIC  
OPTION (M SERIES BI-PHASIC)**Nov 5, 2003  
90 days to decisionK032439 · Product code: **MKJ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k032439/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Aug 7, 2003
Decision date	Nov 5, 2003
Days to decision	90 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Zoll Medical Corp</b>
Location	Woburn, MA, US
Contact	SCOTT AUGUST
510(k) history	33 submissions · 27 cleared · 1993-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k032439/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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