

**K991556 ZOLL M SERIES 12SL ANALYSIS OPTION**Oct 21, 1999  
170 days to decisionK991556 · Product code: **MKJ** · CardiovascularSource: <https://www.510kdatabase.net/k991556/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - ST
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	May 4, 1999
Decision date	Oct 21, 1999
Days to decision	170 days
Third-party review	No

**APPLICANT**

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Company	<b>Zoll Medical Corp</b>
Location	Woburn, MA, US
Contact	PAUL DIAS
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/k991556/>; 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 July 3, 2026