

**K926172 SERIES 1500 DEFIBRILLATOR, MODIFICATION**Feb 25, 1993  
86 days to decisionK926172 · Product code: **MKJ** · CardiovascularSource: <https://www.510kdatabase.net/k926172/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Automated External Defibrillators (non-wearable) (MKJ)
Date received	Dec 1, 1992
Decision date	Feb 25, 1993
Days to decision	86 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Marquette Electronics, Inc.</b>
Location	Walker, MI, US
Contact	GERRY BUSS
510(k) history	82 submissions · 81 cleared · 1980-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k926172/> 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 May 21, 2026