

**K870622 HEART\*AID 1000**Apr 10, 1987  
57 days to decisionK870622 · Product code: **MKJ** · CardiovascularSource: <https://www.510kdatabase.net/k870622/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)                     |
| Submission type       | Traditional  |
| Device classification | Automated External Defibrillators (non-wearable) (MKJ) |
| Date received         | Feb 12, 1987   |
| Decision date         | Apr 10, 1987   |
| Days to decision      | 57 days  |
| Third-party review    | No   |

**APPLICANT**

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|----------------|---|
| Company        | <b>Cardiac Resucitator Corp.</b>        |
| Location       | Mchenry, IL, US                         |
| Contact        | JOHN L KARPOWICZ                        |
| 510(k) history | 17 submissions · 16 cleared · 1981-1988 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k870622/>; 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 June 29, 2026