

**K955923 MEDI-TRACE COMBINATION DEFIBRILLATION & ECG  
ELECTRODE FOR LAERDAL**Oct 24, 1996  
301 days to decisionK955923 · Product code: **MKJ** · Cardiovascular  
Source: <https://www.510kdatabase.net/k955923/>**SUBMISSION DETAILS**

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|                       |  |
|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)                     |
| Submission type       | Traditional  |
| Device classification | Automated External Defibrillators (non-wearable) (MKJ) |
| Date received         | Dec 28, 1995   |
| Decision date         | Oct 24, 1996   |
| Days to decision      | 301 days   |
| Third-party review    | No   |
| Summary / Statement   | Summary  |

**APPLICANT**

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|----------------|---|
| Company        | <b>Graphic Controls Corp.</b>           |
| Location       | Mchenry, IL, US                         |
| Contact        | KATHLEEN SELOVER                        |
| 510(k) history | 55 submissions · 55 cleared · 1977-1998 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k955923/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated May 21, 2026