

**K872972 TEMPORARY PACEMAKER ELECTRODE**Dec 3, 1987  
127 days to decisionK872972 · Product code: **LDF** · Cardiovascular  
Source: <https://www.510kdatabase.net/k872972/>**SUBMISSION DETAILS**

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|                       |                                       |
|-----------------------|---------------------------------------|
| Decision              | Substantially Equivalent (Cleared)    |
| Submission type       | Traditional                           |
| Device classification | Electrode, Pacemaker, Temporary (LDF) |
| Date received         | Jul 29, 1987                          |
| Decision date         | Dec 3, 1987                           |
| Days to decision      | 127 days                              |
| Third-party review    | No                                    |

**APPLICANT**

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|----------------|---|
| Company        | <b>Oscor Medical Corp.</b>              |
| Location       | Washington, DC, US                      |
| Contact        | BRIAN K CORNISH                         |
| 510(k) history | 31 submissions · 30 cleared · 1985-1997 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k872972/>, 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