

K862330 MODIFIED OSCOR PY PACING LEADSAug 8, 1986
50 days to decisionK862330 · Product code: **DTB** · CardiovascularSource: <https://www.510kdatabase.net/k862330/>**SUBMISSION DETAILS**

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|-----------------------|-------------------------------------|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Traditional |
| Device classification | Permanent Pacemaker Electrode (DTB) |
| Date received | Jun 19, 1986 |
| Decision date | Aug 8, 1986 |
| Days to decision | 50 days |
| Third-party review | No |

APPLICANT

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|----------------|---|
| Company | Oscor Medical Corp. |
| Location | Washington, DC, US |
| Contact | DE GRAAD |
| 510(k) history | 31 submissions · 30 cleared · 1985-1997 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k862330/>, Data sourced from FDA 510(k) public records (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. - Generated July 3, 2026