

**K870700 MICRO-GUIDE**Dec 3, 1987  
287 days to decisionK870700 · Product code: **DQO** · CardiovascularSource: <https://www.510kdatabase.net/k870700/>**SUBMISSION DETAILS**

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|-----------------------|---|
| Decision              | Substantially Equivalent (Cleared)        |
| Submission type       | Traditional                               |
| Device classification | Catheter, Intravascular, Diagnostic (DQO) |
| Date received         | Feb 19, 1987                              |
| Decision date         | Dec 3, 1987                               |
| Days to decision      | 287 days                                  |
| Third-party review    | No  |

**APPLICANT**

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|----------------|--|
| Company        | <b>Catheter Research C/O Burditt, Bowles &amp; Radzius</b> |
| Location       | Chicago, IL, US  |
| Contact        | JOSEPH R RADZIUS   |
| 510(k) history | 8 submissions · 8 cleared · 1987-1993                      |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k870700/>, 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 6, 2026