

**K992942 LEMAITRE OCCLUSION CATHETER**Apr 13, 2000  
226 days to decisionK992942 · Product code: **MJN** · CardiovascularSource: <https://www.510kdatabase.net/k992942/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)                 |
| Submission type       | Traditional  |
| Device classification | Catheter, Intravascular Occluding, Temporary (MJN) |
| Date received         | Aug 31, 1999                                       |
| Decision date         | Apr 13, 2000                                       |
| Days to decision      | 226 days   |
| Third-party review    | No   |
| Summary / Statement   | Statement  |

**APPLICANT**

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|----------------|---|
| Company        | <b>Vascutech, Inc.</b>                  |
| Location       | Washington, DC, US                      |
| Contact        | TRENT G KAMKE                           |
| 510(k) history | 14 submissions · 14 cleared · 1985-2000 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k992942/> 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 May 20, 2026