

**K890717 MODIFICATION HEMOFILTRATION CATHETER HF-100**Apr 5, 1989  
51 days to decisionK890717 · Product code: **LFK** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k890717/>**SUBMISSION DETAILS**

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|                       |                                    |
|-----------------------|------------------------------------|
| Decision              | Substantially Equivalent (Cleared) |
| Submission type       | Traditional                        |
| Device classification | Catheter, Femoral (LFK)            |
| Date received         | Feb 13, 1989                       |
| Decision date         | Apr 5, 1989                        |
| Days to decision      | 51 days                            |
| Third-party review    | No                                 |

**APPLICANT**

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|----------------|---|
| Company        | <b>Vas-Cath, Inc.</b>                   |
| Location       | Mississauga, Ontario, CA                |
| Contact        | BETTY-ANN BUTCHER                       |
| 510(k) history | 27 submissions · 27 cleared · 1988-1998 |

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k890717/>; 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 1, 2026