

**K933981 CERVICAL CATHETER**Dec 27, 1994  
498 days to decisionK933981 · Product code: **LKF** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k933981/>**SUBMISSION DETAILS**

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|                       |  |
|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)           |
| Submission type       | Traditional                                  |
| Device classification | Cannula, Manipulator/injector, Uterine (LKF) |
| Date received         | Aug 16, 1993                                 |
| Decision date         | Dec 27, 1994                                 |
| Days to decision      | 498 days                                     |
| Third-party review    | No   |
| Summary / Statement   | Summary                                      |

**APPLICANT**

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|----------------|---|
| Company        | <b>Conceptus, Inc.</b>  |
| Location       | San Carlos, CA, US  |
| Contact        | ALEXIS BALL   |
| Website        | <a href="http://www.conceptus.com">http://www.conceptus.com</a> |
| 510(k) history | 9 submissions · 9 cleared · 1994-1998                           |

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k933981/> 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 June 29, 2026