

**K932993 INTRAUTERINE CATHETER AND INTRODUCER**Jan 6, 1995  
567 days to decisionK932993 · Product code: **MFD** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k932993/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Intrauterine Insemination (MFD)
Date received	Jun 18, 1993
Decision date	Jan 6, 1995
Days to decision	567 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/k932993/> 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