

**K940681 CLEARVIEW UTERINE MANIPULATOR**Jul 26, 1994  
166 days to decisionK940681 · Product code: **LKF** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k940681/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Manipulator/injector, Uterine (LKF)
Date received	Feb 10, 1994
Decision date	Jul 26, 1994
Days to decision	166 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Clinical Innovations, Inc.</b>
Location	Murray, UT, US
Contact	WILLIAM DEAN WALLACE
510(k) history	9 submissions · 9 cleared · 1994-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k940681/> 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