

**K131781 CLEARVIEW TOTAL**May 28, 2014  
345 days to decisionK131781 · Product code: **LKF** · Obstetrics & Gynecology  
Source: <https://www.510kdatabase.net/k131781/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Manipulator/injector, Uterine (LKF)
Date received	Jun 17, 2013
Decision date	May 28, 2014
Days to decision	345 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Clinical Innovations, LLC</b>
Location	Murray, UT, US
Contact	BRIAN MCROBERTS
510(k) history	9 submissions · 9 cleared · 2007-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k131781/> 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 26, 2026