

**K123258 THE ADVANCE CATHETER FOR HSG AND SIS**Jan 31, 2013  
105 days to decisionK123258 · Product code: **LKF** · Obstetrics & Gynecology  
Source: <https://www.510kdatabase.net/k123258/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Cannula, Manipulator/injector, Uterine (LKF)
Date received	Oct 18, 2012
Decision date	Jan 31, 2013
Days to decision	105 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Catheter Research, Inc.</b>
Location	Indianapolis, IN, US
Contact	BABACAR DIOUF
510(k) history	8 submissions · 8 cleared · 1994-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k123258/> 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 25, 2026