

**K931122 OVAMED HSG CATHETER**Jan 30, 1995  
697 days to decisionK931122 · Product code: **LKF** · Obstetrics & Gynecology  
Source: <https://www.510kdatabase.net/k931122/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Manipulator/injector, Uterine (LKF)
Date received	Mar 4, 1993
Decision date	Jan 30, 1995
Days to decision	697 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Ovamed Corp.</b>
Location	Sunnyvale, CA, US
Contact	CHRISTINE DECARIA
510(k) history	5 submissions · 5 cleared · 1995-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k931122/> 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