

**K961752 H/S CATHETER SET(61-3005, 61-3007, 61-3605, 61-3607)**Aug 8, 1996  
94 days to decisionK961752 · Product code: **LKF** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k961752/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Manipulator/injector, Uterine (LKF)
Date received	May 6, 1996
Decision date	Aug 8, 1996
Days to decision	94 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Ackrad Laboratories</b>
Location	Mchenry, IL, US
Contact	BERNARD ACKERMAN
510(k) history	42 submissions · 41 cleared · 1979-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k961752/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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