

**K953034 H/S CATHETER SET(61-3005, 61-3007, 61-3605, 61-3607)**

Aug 29, 1995  
71 days to decision

K953034 · Product code: **LKF** · Obstetrics & Gynecology  
Source: <https://www.510kdatabase.net/k953034/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Manipulator/injector, Uterine (LKF)
Date received	Jun 19, 1995
Decision date	Aug 29, 1995
Days to decision	71 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, PH.D.
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/k953034/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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