

K021272 SIB CATHETER, MODEL 61-7005May 21, 2002
29 days to decisionK021272 · Product code: **LKF** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k021272/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Cannula, Manipulator/injector, Uterine (LKF)
Date received	Apr 22, 2002
Decision date	May 21, 2002
Days to decision	29 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Ackrad Laboratories
Location	Mchenry, IL, US
Contact	RICHARD HETTENBACH
510(k) history	42 submissions · 41 cleared · 1979-2002

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k021272/> Data sourced from FDA 510(k) public records (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. - Generated May 25, 2026