FDA 510(k) Premarket Notification

K021272 SIB CATHETER, MODEL 61-7005

CLEARED

May 21, 2002
29 days to decision

K021272 · Product code: **LKF** · Obstetrics & Gynecology
Source: 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