FDA 510(k) Premarket Notification

K961752 H/S CATHETER SET(61-3005, 61-3007, 61-3605, 61-3607)

CLEARED

Aug 8, 1996
94 days to decision

K961752 · Product code: **LKF** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k961752/

SUBMISSION DETAILS

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

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