FDA 510(k) Premarket Notification

K010056 UTERINE MANIPULATOR INJECTOR CANNULA, STERILE, MODEL R 57-450

CLEARED

Feb 1, 2001
24 days to decision

K010056 · Product code: **LKF** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k010056/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Manipulator/injector, Uterine (LKF)
Date received	Jan 8, 2001
Decision date	Feb 1, 2001
Days to decision	24 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**A & A Medical, Inc.**
Location	Branford, CT, US
Contact	JIHAD MANSOUR
510(k) history	23 submissions · 23 cleared · 1988-2001