

**K010056 UTERINE MANIPULATOR INJECTOR CANNULA,
STERILE, MODEL R 57-450**Feb 1, 2001
24 days to decisionK010056 · Product code: **LKF** · Obstetrics & GynecologySource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k010056/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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