

K772230 CANNULA, PERFUSION, FALLOPIAN TUBEJan 3, 1978
28 days to decisionK772230 · Product code: **LKF** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k772230/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Manipulator/injector, Uterine (LKF)
Date received	Dec 6, 1977
Decision date	Jan 3, 1978
Days to decision	28 days
Third-party review	No

APPLICANT

Company	J. Sklar Mfg. Co., Inc.
Location	Mchenry, IL, US
510(k) history	18 submissions · 18 cleared · 1977-1984

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Device record: <https://www.510kdatabase.net/k772230/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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