

K023051 INTRAUTERINE DELIVERY SYSTEM - INJECTORDec 12, 2002
90 days to decisionK023051 · Product code: **LKF** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k023051/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Manipulator/injector, Uterine (LKF)
Date received	Sep 13, 2002
Decision date	Dec 12, 2002
Days to decision	90 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Prosurg, Inc.
Location	San Jose, CA, US
Contact	ASHVIN DESAI
510(k) history	16 submissions · 16 cleared · 2000-2016

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