

**K020292 UTERINE INJECTOR**Apr 17, 2002  
79 days to decisionK020292 · Product code: **LKF** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k020292/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Manipulator/injector, Uterine (LKF)
Date received	Jan 28, 2002
Decision date	Apr 17, 2002
Days to decision	79 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Catheter Research, Inc.</b>
Location	Indianapolis, IN, US
Contact	JOHN A STEEN
510(k) history	8 submissions · 8 cleared · 1994-2013

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