FDA 510(k) Premarket Notification

K020292 UTERINE INJECTOR

CLEARED

Apr 17, 2002
79 days to decision

K020292 · Product code: **LKF** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k020292/

SUBMISSION DETAILS

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

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