FDA 510(k) Premarket Notification

K122820 VIKY UP

CLEARED

Jun 18, 2013
277 days to decision

K122820 · Product code: **LKF** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k122820/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Manipulator/injector, Uterine (LKF)
Date received	Sep 14, 2012
Decision date	Jun 18, 2013
Days to decision	277 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Endocontrol**
Location	La Tronche, FR
Contact	HUGUEL CARINE
510(k) history	2 submissions · 2 cleared · 2008-2013