FDA 510(k) Premarket Notification

K131781 CLEARVIEW TOTAL

CLEARED

May 28, 2014
345 days to decision

K131781 · Product code: **LKF** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k131781/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Manipulator/injector, Uterine (LKF)
Date received	Jun 17, 2013
Decision date	May 28, 2014
Days to decision	345 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Clinical Innovations, LLC**
Location	Murray, UT, US
Contact	BRIAN MCROBERTS
510(k) history	9 submissions · 9 cleared · 2007-2025