

**K142545 SonoFlow Sonohysterography and Sonohysterosalpingography Device**Mar 20, 2015  
191 days to decisionK142545 · Product code: **LKF** · Obstetrics & Gynecology  
Source: <https://www.510kdatabase.net/k142545/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Manipulator/injector, Uterine (LKF)
Date received	Sep 10, 2014
Decision date	Mar 20, 2015
Days to decision	191 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Crossbay Medical, Inc.</b>
Location	Irvine, CA, US
Contact	Piush Vidyarthi
510(k) history	3 submissions · 3 cleared · 2014-2017

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k142545/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 2, 2026