

**K130758 KRONNER SIDE-KICK UTERINE MANIPULATOR  
HOLDER**Dec 11, 2013  
266 days to decisionK130758 · Product code: **LKF** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k130758/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Manipulator/injector, Uterine (LKF)
Date received	Mar 20, 2013
Decision date	Dec 11, 2013
Days to decision	266 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Kronner Prototypes, Inc.</b>
Location	Roseburg, OR, US
Contact	RICHARD F KRONNER, M.D.
510(k) history	4 submissions · 4 cleared · 1994-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k130758/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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