

**K932394 BERNSTEIN(TM) UTERINE MANIPULATOR**Jan 26, 1995  
618 days to decisionK932394 · Product code: **LKF** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k932394/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Manipulator/injector, Uterine (LKF)
Date received	May 18, 1993
Decision date	Jan 26, 1995
Days to decision	618 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Gm Engineering, Inc.</b>
Location	La Verne, CA, US
Contact	DAVID C STEFFIN
510(k) history	11 submissions · 11 cleared · 1993-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k932394/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 29, 2026