FDA 510(k) Premarket Notification

K932394 BERNSTEIN(TM) UTERINE MANIPULATOR

CLEARED

Jan 26, 1995
618 days to decision

K932394 · Product code: **LKF** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k932394/

SUBMISSION DETAILS

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

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