FDA 510(k) Premarket Notification

K933981 CERVICAL CATHETER

CLEARED

Dec 27, 1994
498 days to decision

K933981 · Product code: **LKF** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k933981/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Manipulator/injector, Uterine (LKF)
Date received	Aug 16, 1993
Decision date	Dec 27, 1994
Days to decision	498 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Conceptus, Inc.**
Location	San Carlos, CA, US
Contact	ALEXIS BALL
Website	http://www.conceptus.com
510(k) history	9 submissions · 9 cleared · 1994-1998