FDA 510(k) Premarket Notification

K931122 OVAMED HSG CATHETER

CLEARED

Jan 30, 1995
697 days to decision

K931122 · Product code: **LKF** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k931122/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Manipulator/injector, Uterine (LKF)
Date received	Mar 4, 1993
Decision date	Jan 30, 1995
Days to decision	697 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Ovamed Corp.**
Location	Sunnyvale, CA, US
Contact	CHRISTINE DECARIA
510(k) history	5 submissions · 5 cleared · 1995-1995