FDA 510(k) Premarket Notification

K932993 INTRAUTERINE CATHETER AND INTRODUCER

CLEARED

Jan 6, 1995
567 days to decision

K932993 · Product code: **MFD** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k932993/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Intrauterine Insemination (MFD)
Date received	Jun 18, 1993
Decision date	Jan 6, 1995
Days to decision	567 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