FDA 510(k) Premarket Notification

K023106 PROMOTOR

CLEARED

Nov 5, 2002
48 days to decision

K023106 · Product code: **MQL** · Obstetrics & Gynecology
Source: https://www.510kdatabase.net/k023106/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Media, Reproductive (MQL)
Date received	Sep 18, 2002
Decision date	Nov 5, 2002
Days to decision	48 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Ceres Fertility, Inc.**
Location	Irvine, CA, US
Contact	GRACE HOLLAND
510(k) history	2 submissions · 2 cleared · 2002-2003