

K023106 PROMOTORNov 5, 2002
48 days to decisionK023106 · Product code: **MQL** · Obstetrics & GynecologySource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k023106/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated June 29, 2026