

K921518 LIFETEK IUI CATHETERMay 10, 1994
770 days to decisionK921518 · Product code: **MFD** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k921518/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Cannula, Intrauterine Insemination (MFD)
Date received	Mar 31, 1992
Decision date	May 10, 1994
Days to decision	770 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Lifetek Medical, Inc.
Location	Deforest, WI, US
Contact	MICHAEL KVALO
510(k) history	2 submissions · 2 cleared · 1994-1999

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k921518/> 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