

K912075 C-SECTION TRAYApr 7, 1992
333 days to decisionK912075 · Product code: **KNA** · Obstetrics & GynecologySource: <https://www.510kdatabase.net/k912075/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent - K
Submission type	Traditional
Device classification	Instrument, Manual, Specialized Obstetric-gynecologic (KNA)
Date received	May 10, 1991
Decision date	Apr 7, 1992
Days to decision	333 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	Medical Device Inspection Co., Inc.
Location	Great Neck, NY, US
Contact	ALAN P.SCHWARTZ
510(k) history	30 submissions · 26 cleared · 1990-1995

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k912075/> 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 30, 2026