FDA 510(k) Premarket Notification

K912075 C-SECTION TRAY

CLEARED

Apr 7, 1992
333 days to decision

K912075 · Product code: **KNA** · Obstetrics & Gynecology
Source: 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