

K912052 HEART CATH TRAYAug 20, 1991
103 days to decisionK912052 · Product code: **FSM** · General & Plastic SurgerySource: <https://www.510kdatabase.net/k912052/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tray, Surgical, Instrument (FSM)
Date received	May 9, 1991
Decision date	Aug 20, 1991
Days to decision	103 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/k912052/> 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