

K893233 DISPOSABLE NEEDLE, ASPIRATION AND INJECTIONJan 4, 1990
251 days to decisionK893233 · Product code: **FMF** · General Hospital
Source: <https://www.510kdatabase.net/k893233/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Apr 28, 1989
Decision date	Jan 4, 1990
Days to decision	251 days
Third-party review	No

APPLICANT

Company	Pioneer Intl. Co.
Location	Hidden Hills, CA, US
Contact	DAVID HSIUNG
510(k) history	1 submissions · 1 cleared · 1990-1990

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k893233/>; 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 July 8, 2026