

K954189 EASI-JECTOR CONTROL SYSTEMOct 19, 1995
43 days to decisionK954189 · Product code: **FMF** · General Hospital
Source: <https://www.510kdatabase.net/k954189/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Sep 6, 1995
Decision date	Oct 19, 1995
Days to decision	43 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	North, Inc.
Location	Northfield, MN, US
Contact	JIM POKORNEY
510(k) history	2 submissions · 2 cleared · 1995-1997

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k954189/> 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 6, 2026