

K820164 EMPTY STERILE CARPUJECTFeb 24, 1982
35 days to decisionK820164 · Product code: **FMF** · General HospitalSource: <https://www.510kdatabase.net/k820164/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Jan 20, 1982
Decision date	Feb 24, 1982
Days to decision	35 days
Third-party review	No

APPLICANT

Company	Sterling Drug, Inc.
Location	Mchenry, IL, US
510(k) history	9 submissions · 9 cleared · 1978-1990

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k820164/>; 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