FDA 510(k) Premarket Notification

K820164 EMPTY STERILE CARPUJECT

CLEARED

Feb 24, 1982
35 days to decision

K820164 · Product code: **FMF** · General Hospital
Source: 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