FDA 510(k) Premarket Notification

K812888 SYRINGE, PISTON

CLEARED

Nov 10, 1981
26 days to decision

K812888 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k812888/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Oct 15, 1981
Decision date	Nov 10, 1981
Days to decision	26 days
Third-party review	No

APPLICANT

Company	**Frisco Products, Inc.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1981-1982