FDA 510(k) Premarket Notification

K833460 SYRINGE,PISTON NEEDLE, HOLLOW LUMEN

CLEARED

Nov 28, 1983
54 days to decision

K833460 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k833460/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Oct 5, 1983
Decision date	Nov 28, 1983
Days to decision	54 days
Third-party review	No

APPLICANT

Company	**Aldrich Chemical Co., Inc.**
Location	Mchenry, IL, US
510(k) history	2 submissions · 2 cleared · 1983-1983