FDA 510(k) Premarket Notification

K770425 NEEDLES, STERILE, AIRWAY

CLEARED

Apr 15, 1977
39 days to decision

K770425 · Product code: **FMI** · General Hospital
Source: https://www.510kdatabase.net/k770425/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Needle, Hypodermic, Single Lumen (FMI)
Date received	Mar 7, 1977
Decision date	Apr 15, 1977
Days to decision	39 days
Third-party review	No

APPLICANT

Company	**Pfizer Pharmaceuticals**
Location	Mchenry, IL, US
510(k) history	6 submissions · 5 cleared · 1977-1989