FDA 510(k) Premarket Notification

K801430 CONTROL SYRINGE

CLEARED

Jul 8, 1980
21 days to decision

K801430 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k801430/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Jun 17, 1980
Decision date	Jul 8, 1980
Days to decision	21 days
Third-party review	No

APPLICANT

Company	**Abco Dealers, Inc.**
Location	Mchenry, IL, US
510(k) history	127 submissions · 127 cleared · 1976-1991