FDA 510(k) Premarket Notification

K820384 GLASS SYRINGES, DRUG, LUER LOCK

CLEARED

Apr 15, 1982
65 days to decision

K820384 · Product code: **FMF** · General Hospital
Source: https://www.510kdatabase.net/k820384/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Syringe, Piston (FMF)
Date received	Feb 9, 1982
Decision date	Apr 15, 1982
Days to decision	65 days
Third-party review	No

APPLICANT

Company	**Procedure Products, Inc.**
Location	Mchenry, IL, US
510(k) history	16 submissions · 16 cleared · 1981-2017