FDA 510(k) Premarket Notification

K810921 MALE LUER LOCK PLUGS

CLEARED

May 21, 1981
45 days to decision

K810921 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k810921/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Apr 6, 1981
Decision date	May 21, 1981
Days to decision	45 days
Third-party review	No

APPLICANT

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