FDA 510(k) Premarket Notification

K811110 MODELS: 52-01-0, 52-01-1

CLEARED

Jun 11, 1981
49 days to decision

K811110 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k811110/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Apr 23, 1981
Decision date	Jun 11, 1981
Days to decision	49 days
Third-party review	No

APPLICANT

Company	**Y**
Location	Mchenry, IL, US
510(k) history	78 submissions · 78 cleared · 1978-1996