FDA 510(k) Premarket Notification

K901949 IV SET

CLEARED

Sep 12, 1990
135 days to decision

K901949 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k901949/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Apr 30, 1990
Decision date	Sep 12, 1990
Days to decision	135 days
Third-party review	No

APPLICANT

Company	**Jfv, Inc.**
Location	Spencer, IN, US
Contact	VANCE, SR.
510(k) history	1 submissions · 1 cleared · 1990-1990