FDA 510(k) Premarket Notification

K901151 I.V. ADMINISTRATION SET CONNECTING DEVICE

CLEARED

Jul 2, 1990
112 days to decision

K901151 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k901151/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Mar 12, 1990
Decision date	Jul 2, 1990
Days to decision	112 days
Third-party review	No

APPLICANT

Company	**Dmd, Div. of Casta Pacific Co.**
Location	Nevada City, CA, US
Contact	JOHN T WRIGHT
510(k) history	1 submissions · 1 cleared · 1990-1990