FDA 510(k) Premarket Notification

K901081 PRO-LOCK(TM) SHIELDING NEEDLE CONNECTOR ASSEMBLY

CLEARED

Jun 8, 1990
93 days to decision

K901081 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k901081/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Mar 7, 1990
Decision date	Jun 8, 1990
Days to decision	93 days
Third-party review	No

APPLICANT

Company	**Vlv Associates, Inc.**
Location	East Hanover, NJ, US
Contact	VINCENT VAILLANCOURT
510(k) history	12 submissions · 11 cleared · 1986-1997