FDA 510(k) Premarket Notification

K881975 SET, INTRAVASCULAR

CLEARED

Jul 29, 1988
79 days to decision

K881975 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k881975/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	May 11, 1988
Decision date	Jul 29, 1988
Days to decision	79 days
Third-party review	No

APPLICANT

Company	**Primrose Medical, Inc.**
Location	Mchenry, IL, US
Contact	FLETCHER LONGLEY
510(k) history	14 submissions · 14 cleared · 1984-1989