FDA 510(k) Premarket Notification

K900336 V SET

CLEARED

Oct 4, 1990
254 days to decision

K900336 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k900336/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jan 23, 1990
Decision date	Oct 4, 1990
Days to decision	254 days
Third-party review	No

APPLICANT

Company	**Go Medical Industries Pty. , Ltd.**
Location	Walker, MI, US
Contact	DAVID CAPES
510(k) history	11 submissions · 11 cleared · 1984-2001