FDA 510(k) Premarket Notification

K885030 STANDARD ADMINISTRATION SETS

CLEARED

Jan 24, 1989
50 days to decision

K885030 · Product code: **FPA** · General Hospital
Source: https://www.510kdatabase.net/k885030/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Dec 5, 1988
Decision date	Jan 24, 1989
Days to decision	50 days
Third-party review	No
Combination product	No
PCCP authorized	No

APPLICANT

Company	**3M Company**
Location	White City, OR, US
Contact	J SUEDKAMP
Website	http://www.3m.com/
510(k) history	331 submissions · 322 cleared · 1976-2025