FDA 510(k) Premarket Notification

K891005 INTRAVASCULAR ADMINI. SET MODELS 60030 & 60050

CLEARED

May 11, 1989
72 days to decision

K891005 · Product code: **FPK** · General Hospital
Source: https://www.510kdatabase.net/k891005/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Tubing, Fluid Delivery (FPK)
Date received	Feb 28, 1989
Decision date	May 11, 1989
Days to decision	72 days
Third-party review	No
Combination product	No
PCCP authorized	No

APPLICANT

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