

K885030 STANDARD ADMINISTRATION SETSJan 24, 1989
50 days to decisionK885030 · Product code: **FPA** · General HospitalSource: <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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k885030/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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