

K904217 PRO-LOK (TM) EXTENSION SETSOct 3, 1990
21 days to decisionK904217 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k904217/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Sep 12, 1990
Decision date	Oct 3, 1990
Days to decision	21 days
Third-party review	No

APPLICANT

Company	Viv Associates, Inc.
Location	East Hanover, NJ, US
Contact	VINCENT VAILLANCOURT
510(k) history	12 submissions · 11 cleared · 1986-1997

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k904217/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 6, 2026