

K900909 IN-LINE VIAL CONNECTION SITEMay 18, 1990
80 days to decisionK900909 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k900909/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Feb 27, 1990
Decision date	May 18, 1990
Days to decision	80 days
Third-party review	No

APPLICANT

Company	Baxter Healthcare Corp
Location	Mchenry, IL, US
Contact	PATRICIA S BARSANTI
510(k) history	505 submissions · 496 cleared · 1977-2019

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k900909/>; 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 May 17, 2026