

K915390 3 LEAD EXTENSION WITH 2 INTERLINK INJECTION SITESMar 12, 1992
104 days to decisionK915390 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k915390/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Nov 29, 1991
Decision date	Mar 12, 1992
Days to decision	104 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k915390/> 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