

K891176 INJECTION SITE PLUG (INTRAVASCULAR ADMINISTRATION)May 18, 1989
73 days to decisionK891176 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k891176/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Mar 6, 1989
Decision date	May 18, 1989
Days to decision	73 days
Third-party review	No

APPLICANT

Company	Multi-Med, Inc.
Location	West Swanzey, NH, US
Contact	ALAN P REID
510(k) history	7 submissions · 7 cleared · 1989-1989

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k891176/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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