

K903541 SAFETY INJECTION PORTSep 14, 1990
38 days to decisionK903541 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k903541/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Aug 7, 1990
Decision date	Sep 14, 1990
Days to decision	38 days
Third-party review	No

APPLICANT

Company	Du-Call Miller Plastics, Inc.
Location	Lisle, IL, US
Contact	W. C MILLER
510(k) history	1 submissions · 1 cleared · 1990-1990

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