

K930751 DURHAMSep 3, 1993
203 days to decisionK930751 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k930751/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Feb 12, 1993
Decision date	Sep 3, 1993
Days to decision	203 days
Third-party review	No
Summary / Statement	Statement
Other names	EXTENSION TUBE LUER LOCK W/INJECTION SITE

APPLICANT

Company	Intl. Medical Consultants
Location	Washington Crossing, PA, US
Contact	JOHN ROMANO
510(k) history	6 submissions · 6 cleared · 1993-1993

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k930751/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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