

**K001233 ACTI-FLEX FLUID DELIVERY SYSTEM OR ACTI-FLEX
F D S**Dec 13, 2000
240 days to decisionK001233 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k001233/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Apr 17, 2000
Decision date	Dec 13, 2000
Days to decision	240 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	R-Group Intl.
Location	Gainesville, FL, US
Contact	KERRY ANNE KAHN
510(k) history	7 submissions · 7 cleared · 1992-2000

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k001233/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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