

K982123 CONTRAST CONTROL DEVICEJul 21, 1998
35 days to decisionK982123 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k982123/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jun 16, 1998
Decision date	Jul 21, 1998
Days to decision	35 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Medex, Inc.
Location	Mchenry, IL, US
Contact	CINDY CHENETSKI
510(k) history	48 submissions · 46 cleared · 1977-2005

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