

K841859 SECURA SETJul 3, 1984
60 days to decisionK841859 · Product code: **FPA** · General Hospital
Source: <https://www.510kdatabase.net/k841859/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	May 4, 1984
Decision date	Jul 3, 1984
Days to decision	60 days
Third-party review	No

APPLICANT

Company	Diatek, Inc.
Location	Mchenry, IL, US
510(k) history	15 submissions · 15 cleared · 1979-2002

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Device record: <https://www.510kdatabase.net/k841859/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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