

**K830377 SECURA SET PRIMARY ADMIN. SET**Mar 1, 1983  
25 days to decisionK830377 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k830377/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Feb 4, 1983
Decision date	Mar 1, 1983
Days to decision	25 days
Third-party review	No

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k830377/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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