

**K881975 SET, INTRAVASCULAR**Jul 29, 1988  
79 days to decisionK881975 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k881975/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	May 11, 1988
Decision date	Jul 29, 1988
Days to decision	79 days
Third-party review	No

**APPLICANT**

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Company	<b>Primrose Medical, Inc.</b>
Location	Mchenry, IL, US
Contact	FLETCHER LONGLEY
510(k) history	14 submissions · 14 cleared · 1984-1989

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k881975/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 27, 2026