

**K900336 V SET**Oct 4, 1990  
254 days to decisionK900336 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k900336/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Jan 23, 1990
Decision date	Oct 4, 1990
Days to decision	254 days
Third-party review	No

**APPLICANT**

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Company	<b>Go Medical Industries Pty. , Ltd.</b>
Location	Walker, MI, US
Contact	DAVID CAPES
510(k) history	11 submissions · 11 cleared · 1984-2001

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k900336/>; 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 June 30, 2026