

**K020419 V SET**Aug 29, 2003  
568 days to decisionK020419 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k020419/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Feb 7, 2002
Decision date	Aug 29, 2003
Days to decision	568 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>P.T. Greenleaf</b>
Location	Perth, AU
Contact	GEORGE O'NEIL
510(k) history	2 submissions · 2 cleared · 2003-2003

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k020419/> 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 July 2, 2026