

**K921391 SOLUTION ADMINISTRATION SET**Sep 2, 1992  
163 days to decisionK921391 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k921391/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Mar 23, 1992
Decision date	Sep 2, 1992
Days to decision	163 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Douglas Medical Products Corp.</b>
Location	Mundelein, IL, US
Contact	DOUGLAS JOHNSON
510(k) history	19 submissions · 19 cleared · 1992-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k921391/> 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 22, 2026