

**K955732 VENTED SOLUTION SET**Feb 21, 1996  
64 days to decisionK955732 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k955732/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Dec 19, 1995
Decision date	Feb 21, 1996
Days to decision	64 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Distrionic Sterile Products, Inc.</b>
Location	Portsmouth, NH, US
Contact	JOYCE MCDUGAL
510(k) history	8 submissions · 8 cleared · 1995-1996

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k955732/> 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 6, 2026