

**K943770 LOW VOLUME MULTIPORT**Apr 10, 1995  
251 days to decisionK943770 · Product code: **FPA** · General HospitalSource: <https://www.510kdatabase.net/k943770/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Aug 2, 1994
Decision date	Apr 10, 1995
Days to decision	251 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>B. Braun of America, Inc.</b>
Location	Bethlehem, PA, US
Contact	MARK S ALSBERGE
510(k) history	19 submissions · 16 cleared · 1991-2000

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k943770/> 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 21, 2026