

**K912069 AVI MODEL 4225 VENTED PIGGYBACK IV ADMIN SET**Oct 14, 1993  
888 days to decisionK912069 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k912069/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	May 10, 1991
Decision date	Oct 14, 1993
Days to decision	888 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>3M Avi, Inc.</b>
Location	St. Paul, MN, US
Contact	VON BUSCH
510(k) history	18 submissions · 18 cleared · 1990-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k912069/> 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 7, 2026