

**K896161 AVI MODEL 2012, 1.2 MICRON FILTER IV  
ADMINISTRA.**Jan 11, 1990  
79 days to decisionK896161 · Product code: **FPA** · General Hospital  
Source: <https://www.510kdatabase.net/k896161/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Set, Administration, Intravascular (FPA)
Date received	Oct 24, 1989
Decision date	Jan 11, 1990
Days to decision	79 days
Third-party review	No
Combination product	No
PCCP authorized	No

**APPLICANT**

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Company	<b>3M Company</b>
Location	White City, OR, US
Contact	VON BUSCH
Website	<a href="http://www.3m.com/">http://www.3m.com/</a>
510(k) history	331 submissions · 322 cleared · 1976-2025

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k896161/>. Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated June 28, 2026