

**K910502 VESSEL DILATORS**Apr 29, 1991  
83 days to decisionK910502 · Product code: **FKA** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k910502/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dilator, Vessel (FKA)
Date received	Feb 5, 1991
Decision date	Apr 29, 1991
Days to decision	83 days
Third-party review	No

**APPLICANT**

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Company	<b>Akcess Medical Products, Inc.</b>
Location	New Brunswick, NJ, US
Contact	BALBIR B KAPANY
510(k) history	36 submissions · 28 cleared · 1989-1994

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k910502/>; 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 June 30, 2026