

**K890203 ADMS BIOPSY NEEDLE**Mar 8, 1989  
50 days to decisionK890203 · Product code: **FCG** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k890203/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Biopsy Needle (FCG)
Date received	Jan 17, 1989
Decision date	Mar 8, 1989
Days to decision	50 days
Third-party review	No

**APPLICANT**

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Company	<b>Advanced Diagnostic Medical Systems, Inc.</b>
Location	Dublin, CA, US
Contact	GARY J ALLSEBROOK
510(k) history	4 submissions · 4 cleared · 1988-1989

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k890203/>; 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