

**K902788 CLEM TM**Sep 12, 1990  
78 days to decisionK902788 · Product code: **FIB** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k902788/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Protector, Transducer, Dialysis (FIB)
Date received	Jun 26, 1990
Decision date	Sep 12, 1990
Days to decision	78 days
Third-party review	No

**APPLICANT**

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Company	<b>Arbo Medical, Inc.</b>
Location	Wilton, CT, US
Contact	DAWN MOORE
510(k) history	15 submissions · 15 cleared · 1989-1995

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k902788/>; 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 2, 2026