

**K863823 AMICON DIAFILTER HEMOFILTER**Nov 6, 1986  
37 days to decisionK863823 · Product code: **KDI** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k863823/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Sep 30, 1986
Decision date	Nov 6, 1986
Days to decision	37 days
Third-party review	No

**APPLICANT**

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Company	<b>Amicon, Inc.</b>
Location	Mchenry, IL, US
Contact	JAMES M DELANEY
510(k) history	20 submissions · 20 cleared · 1976-1993

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k863823/>; 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 14, 2026