

**K812353 KF-E SERIES OF HOLLOW FIBER DIALYZERS**Sep 16, 1981  
28 days to decisionK812353 · Product code: **FJI** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k812353/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, Capillary, Hollow Fiber (FJI)
Date received	Aug 19, 1981
Decision date	Sep 16, 1981
Days to decision	28 days
Third-party review	No

**APPLICANT**

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Company	<b>Medisystems Corp.</b>
Location	Mchenry, IL, US
510(k) history	22 submissions · 22 cleared · 1981-2007

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k812353/> 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 29, 2026