

**K852251 FRESENIUS HEMOFLOW F60 & FRESENIUS HEMOFLOW F40**Jul 25, 1985  
63 days to decisionK852251 · Product code: **KDI** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k852251/>**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	May 23, 1985
Decision date	Jul 25, 1985
Days to decision	63 days
Third-party review	No

**APPLICANT**

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Company	<b>Seratronics, Inc.</b>
Location	Mchenry, IL, US
Contact	LIPPS, PH.D.
510(k) history	21 submissions · 21 cleared · 1982-1996

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k852251/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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