

**K870724 FRESENIUS HEMOFLOW F60 AND F80**Apr 1, 1987  
36 days to decisionK870724 · Product code: **KDI** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k870724/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Feb 24, 1987
Decision date	Apr 1, 1987
Days to decision	36 days
Third-party review	No

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k870724/>; 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