FDA 510(k) Premarket Notification

K870724 FRESENIUS HEMOFLOW F60 AND F80

CLEARED

Apr 1, 1987
36 days to decision

K870724 · Product code: **KDI** · Gastroenterology & Urology
Source: 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	**Seratronics, Inc.**
Location	Mchenry, IL, US
Contact	THOMAS E CANE
510(k) history	21 submissions · 21 cleared · 1982-1996