FDA 510(k) Premarket Notification

K852251 FRESENIUS HEMOFLOW F60 & FRESENIUS HEMOFLOW F40

CLEARED

Jul 25, 1985
63 days to decision

K852251 · Product code: **KDI** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k852251/

SUBMISSION DETAILS

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

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