FDA 510(k) Premarket Notification

K810612 HEMOPHRON

CLEARED

Apr 29, 1981
54 days to decision

K810612 · Product code: **FKQ** · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k810612/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Dialysate Delivery, Central Multiple Patient (FKQ)
Date received	Mar 6, 1981
Decision date	Apr 29, 1981
Days to decision	54 days
Third-party review	No

APPLICANT

Company	**Renal Devices, Inc.**
Location	Mchenry, IL, US
510(k) history	4 submissions · 4 cleared · 1979-1981