

K810612 HEMOPHRONApr 29, 1981
54 days to decisionK810612 · Product code: **FKQ** · Gastroenterology & UrologySource: <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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k810612/> Data sourced from FDA 510(k) public records (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. - Generated July 4, 2026