

K910270 PREMIXED DIALYSATE FOR HEMODIAFILTRATIONApr 18, 1991
85 days to decisionK910270 · Product code: **KPO** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k910270/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialysate Concentrate For Hemodialysis (liquid Or Powder) (KPO)
Date received	Jan 23, 1991
Decision date	Apr 18, 1991
Days to decision	85 days
Third-party review	No

APPLICANT

Company	Baxter Healthcare Corp
Location	Mchenry, IL, US
Contact	BONNIE J SMITH
510(k) history	505 submissions · 496 cleared · 1977-2019

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k910270/>, 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 May 17, 2026