

K923139 HEMOCOR HP HEMOCONCENTRATOR SERIESFeb 18, 1994
599 days to decisionK923139 · Product code: **KDI** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k923139/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Jun 29, 1992
Decision date	Feb 18, 1994
Days to decision	599 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Minntech Corp.
Location	Minneapolis, MN, US
Contact	LEROY J FISCHBACH
510(k) history	33 submissions · 33 cleared · 1987-2012

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k923139/> 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 June 14, 2026