

**K923727 PRIMUS HOLLOW FIBER DIALYZER MOD. 1000, 1350,
2000**Jul 29, 1994
732 days to decisionK923727 · Product code: **KDI** · Gastroenterology & Urology
Source: <https://www.510kdatabase.net/k923727/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Jul 27, 1992
Decision date	Jul 29, 1994
Days to decision	732 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k923727/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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