

**K980859 HEMOCOR HPH MINI HEMOCONCENTRATOR**May 12, 1998  
68 days to decisionK980859 · Product code: **KDI** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k980859/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Mar 5, 1998
Decision date	May 12, 1998
Days to decision	68 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Minntech Corp.</b>
Location	Minneapolis, MN, US
Contact	MARK MURPHY
510(k) history	33 submissions · 33 cleared · 1987-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k980859/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated June 13, 2026