

**K872456 HEMOCOR PLUS HEMOCONCENTRATOR**Sep 15, 1987  
84 days to decisionK872456 · Product code: **KDI** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k872456/>**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	Jun 23, 1987
Decision date	Sep 15, 1987
Days to decision	84 days
Third-party review	No

**APPLICANT**

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Company	<b>Minntech Corp.</b>
Location	Minneapolis, MN, US
Contact	LEROY J FISCHBACH
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/k872456/>; 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