

**K883307 CARDIOFLOW 100  
HEMOCONCENTRATOR/PREPARATION KIT**Oct 12, 1988  
69 days to decisionK883307 · Product code: **KDI** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k883307/>**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	Aug 4, 1988
Decision date	Oct 12, 1988
Days to decision	69 days
Third-party review	No

**APPLICANT**

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Company	<b>Hospal Medical Corp.</b>
Location	Mchenry, IL, US
Contact	CATHERINA MADORMO
510(k) history	55 submissions · 55 cleared · 1977-1989

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k883307/>; 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 May 14, 2026