

**K884365 MULTIFLOW(TM) 60 KIT**Dec 8, 1988  
51 days to decisionK884365 · Product code: **KDI** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k884365/>**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	Oct 18, 1988
Decision date	Dec 8, 1988
Days to decision	51 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/k884365/>; 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