

**K823589 H 12 07**Dec 22, 1982  
16 days to decisionK823589 · Product code: **KDI** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k823589/>**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	Dec 6, 1982
Decision date	Dec 22, 1982
Days to decision	16 days
Third-party review	No

**APPLICANT**

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Company	<b>Hospal Medical Corp.</b>
Location	Mchenry, IL, US
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/k823589/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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