

K812575 MODEL H12-10Oct 19, 1981
40 days to decisionK812575 · Product code: **KDI** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k812575/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Sep 9, 1981
Decision date	Oct 19, 1981
Days to decision	40 days
Third-party review	No

APPLICANT

Company	Hospal Medical Corp.
Location	Mchenry, IL, US
510(k) history	55 submissions · 55 cleared · 1977-1989

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Device record: <https://www.510kdatabase.net/k812575/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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