

K830595 MONITRALApr 28, 1983
63 days to decisionK830595 · Product code: **KDI** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k830595/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Feb 24, 1983
Decision date	Apr 28, 1983
Days to decision	63 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k830595/> Data sourced from FDA 510(k) public records (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. - Generated May 14, 2026