

K840680 CENTRY 2000 DIALYSIS CONTROL UNITApr 20, 1984
65 days to decisionK840680 · Product code: **KDI** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k840680/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Feb 15, 1984
Decision date	Apr 20, 1984
Days to decision	65 days
Third-party review	No

APPLICANT

Company	Cobe Laboratories, Inc.
Location	Mchenry, IL, US
Website	https://www.gambro.com
510(k) history	77 submissions · 77 cleared · 1976-1993

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k840680/>, 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 26, 2026