FDA 510(k) Premarket Notification

K840680 CENTRY 2000 DIALYSIS CONTROL UNIT

CLEARED

Apr 20, 1984
65 days to decision

K840680 · Product code: **KDI** · Gastroenterology & Urology
Source: 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