FDA 510(k) Premarket Notification

K820125 COBE CENTRY 2000 RX

CLEARED

Mar 23, 1982
64 days to decision

K820125 · Gastroenterology & Urology
Source: https://www.510kdatabase.net/k820125/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Date received	Jan 18, 1982
Decision date	Mar 23, 1982
Days to decision	64 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.net
Device record: https://www.510kdatabase.net/k820125/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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