

K931720 LUNDIA ALPHA 700 HEMODIALYZERJul 7, 1994
456 days to decisionK931720 · Product code: **KDI** · Gastroenterology & UrologySource: <https://www.510kdatabase.net/k931720/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Dialyzer, High Permeability With Or Without Sealed Dialysate System (KDI)
Date received	Apr 7, 1993
Decision date	Jul 7, 1994
Days to decision	456 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	C.G.H. Medical, Inc.
Location	Lakewood, CO, US
Contact	VERA BUFFALOE
510(k) history	11 submissions · 11 cleared · 1991-1995

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k931720/> 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