

K904980 DPX-L AND DPX-ALPHA, MODIFICATIONDec 5, 1990
33 days to decisionK904980 · Product code: **KGI** · Radiology
Source: <https://www.510kdatabase.net/k904980/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Densitometer, Bone (KGI)
Date received	Nov 2, 1990
Decision date	Dec 5, 1990
Days to decision	33 days
Third-party review	No

APPLICANT

Company	Lunar
Location	Madison, WI, US
Contact	RICHARD B MAZESS
510(k) history	6 submissions · 6 cleared · 1989-1992

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k904980/>, 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 21, 2026