

K030962 DPX SERIES BRAVO DUO BONE DENSITOMETERJul 25, 2003
120 days to decisionK030962 · Product code: **KGI** · Radiology
Source: <https://www.510kdatabase.net/k030962/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Densitometer, Bone (KGI)
Date received	Mar 27, 2003
Decision date	Jul 25, 2003
Days to decision	120 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Ge Lunar Corp.
Location	Madison, WI, US
Contact	JAMES P RASKOB
510(k) history	3 submissions · 3 cleared · 2001-2003

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k030962/> 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 18, 2026