FDA 510(k) Premarket Notification

K030962 DPX SERIES BRAVO DUO BONE DENSITOMETER

CLEARED

Jul 25, 2003
120 days to decision

K030962 · 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