

K964263 EXPERT-KL AND HAND ACQUISITION AND ANALYSIS SOFTWARE

Nov 7, 1996
13 days to decision

K964263 · Product code: **KGI** · Radiology
Source: <https://www.510kdatabase.net/k964263/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Densitometer, Bone (KGI)
Date received	Oct 25, 1996
Decision date	Nov 7, 1996
Days to decision	13 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Lunar Corp.
Location	Madison, WI, US
Contact	KENNETH D BUOKER
510(k) history	23 submissions · 23 cleared · 1995-2000

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Device record: <https://www.510kdatabase.net/k964263/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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