

K072439 GE LUNAR DPX & PRODIGY SERIES DEXA BONE DENSITOMETERS WITH MOBILE KIT

Sep 14, 2007
15 days to decision

K072439 · Product code: **KGI** · Radiology
Source: <https://www.510kdatabase.net/k072439/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Densitometer, Bone (KGI)
Date received	Aug 30, 2007
Decision date	Sep 14, 2007
Days to decision	15 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Ge Medical Systems Ultrasound and Primary Care Dia
Location	Milwaukee, WI, US
Contact	JAMES P RASKOB
510(k) history	7 submissions · 7 cleared · 2004-2011

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k072439/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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