

**K993176 MULTI-RACIAL REFERENCE POPULATION FOR
PDEXA AND DISCOVERY**Jun 20, 2000
272 days to decisionK993176 · Product code: **KGI** · Radiology
Source: <https://www.510kdatabase.net/k993176/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Densitometer, Bone (KGI)
Date received	Sep 22, 1999
Decision date	Jun 20, 2000
Days to decision	272 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Norland Medical Systems, Inc.
Location	Fort Atkinson, WI, US
Contact	TERRY SCHWALENBERG
510(k) history	3 submissions · 3 cleared · 1999-2000

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k993176/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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