

**K913078 LATERAL SPINE OPT. XR-SERIES X-RAY BONE DENSITOMET**

Sep 24, 1991  
75 days to decision

K913078 · Product code: **KGI** · Radiology  
Source: <https://www.510kdatabase.net/k913078/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Densitometer, Bone (KGI)
Date received	Jul 11, 1991
Decision date	Sep 24, 1991
Days to decision	75 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Norland Corp.</b>
Location	Mchenry, IL, US
Contact	W SCHWALENBERG
510(k) history	15 submissions · 15 cleared · 1979-1998

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k913078/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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