FDA 510(k) Premarket Notification

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

CLEARED

Sep 24, 1991
75 days to decision

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

SUBMISSION DETAILS

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

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