FDA 510(k) Premarket Notification

K904980 DPX-L AND DPX-ALPHA, MODIFICATION

CLEARED

Dec 5, 1990
33 days to decision

K904980 · Product code: **KGI** · Radiology
Source: https://www.510kdatabase.net/k904980/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Densitometer, Bone (KGI)
Date received	Nov 2, 1990
Decision date	Dec 5, 1990
Days to decision	33 days
Third-party review	No

APPLICANT

Company	**Lunar**
Location	Madison, WI, US
Contact	RICHARD B MAZESS
510(k) history	6 submissions · 6 cleared · 1989-1992