FDA 510(k) Premarket Notification

K923823 3D IMAGING

CLEARED

Oct 30, 1992
92 days to decision

K923823 · Product code: **LNH** · Radiology
Source: https://www.510kdatabase.net/k923823/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Jul 30, 1992
Decision date	Oct 30, 1992
Days to decision	92 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

Company	**Resonex, Inc.**
Location	Sunnyvale, CA, US
Contact	BRUCE FLOYD
510(k) history	21 submissions · 21 cleared · 1988-1994