FDA 510(k) Premarket Notification

K930960 ACCESS MAGNETIC RESONANCE DEVICE ACCESSORY

CLEARED

Nov 1, 1993
250 days to decision

K930960 · Product code: **LNH** · Radiology
Source: https://www.510kdatabase.net/k930960/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Feb 24, 1993
Decision date	Nov 1, 1993
Days to decision	250 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Toshiba America Mri, Inc.**
Location	South San Francisco, CA, US
Contact	RON YOKOTA
510(k) history	68 submissions · 68 cleared · 1990-2000