FDA 510(k) Premarket Notification

K926171 MRT-150A, QD SPINE COIL

CLEARED

Apr 27, 1993
140 days to decision

K926171 · Product code: **LNH** · Radiology
Source: https://www.510kdatabase.net/k926171/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Dec 8, 1992
Decision date	Apr 27, 1993
Days to decision	140 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