FDA 510(k) Premarket Notification

K920096 MRT-150 QD KNEE COIL

CLEARED

Jul 10, 1992
184 days to decision

K920096 · Product code: **LNH** · Radiology
Source: https://www.510kdatabase.net/k920096/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Jan 8, 1992
Decision date	Jul 10, 1992
Days to decision	184 days
Third-party review	No
Summary / Statement	Statement

APPLICANT

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