FDA 510(k) Premarket Notification

K910220 MAGNETIC RESONANCE DEVICE ACCESSORY, SM PARTS COIL

CLEARED

Mar 25, 1991
67 days to decision

K910220 · Product code: **LNH** · Radiology
Source: https://www.510kdatabase.net/k910220/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Jan 17, 1991
Decision date	Mar 25, 1991
Days to decision	67 days
Third-party review	No

APPLICANT

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