

**K910220 MAGNETIC RESONANCE DEVICE ACCESSORY, SM
PARTS COIL**Mar 25, 1991
67 days to decisionK910220 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k910220/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 19, 2026