

**K926171 MRT-150A, QD SPINE COIL**Apr 27, 1993  
140 days to decisionK926171 · Product code: **LNH** · Radiology  
Source: <https://www.510kdatabase.net/k926171/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k926171/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 19, 2026