

**K920096 MRT-150 QD KNEE COIL**Jul 10, 1992  
184 days to decisionK920096 · Product code: **LNH** · Radiology  
Source: <https://www.510kdatabase.net/k920096/>**SUBMISSION DETAILS**

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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**

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Company	<b>Toshiba America Mri, Inc.</b>
Location	South San Francisco, CA, US
Contact	BEN KHOSRAVI
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/k920096/> 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