

**K903319 MAGNETIC RESONANCE DIAGNOSTIC DEVICE**Aug 17, 1990  
23 days to decisionK903319 · Product code: **LNH** · Radiology  
Source: <https://www.510kdatabase.net/k903319/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Jul 25, 1990
Decision date	Aug 17, 1990
Days to decision	23 days
Third-party review	No

**APPLICANT**

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Company	<b>GE Medical Systems</b>
Location	Milwaukee, WI, US
Contact	LARRY A KROGER,
510(k) history	169 submissions · 166 cleared · 1989-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k903319/>; 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 27, 2026