

**K924426 MAGNETIC RESONANCE DIAG DEVICE ACCESS  
APPLIC PKG**Dec 18, 1992  
108 days to decisionK924426 · Product code: LNH · Radiology  
Source: <https://www.510kdatabase.net/k924426/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Sep 1, 1992
Decision date	Dec 18, 1992
Days to decision	108 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Siemens Medical Solutions USA, Inc.</b>
Location	Hoffman Estates, IL, US
Contact	KATHLEEN RUTHERFORD
510(k) history	778 submissions · 778 cleared · 1980-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k924426/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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