

**K900889 MAGNETIC RESONANCE DIAGNOSTIC DEVICE  
ACCESSORIES**Mar 23, 1990  
24 days to decisionK900889 · Product code: LNH · Radiology  
Source: <https://www.510kdatabase.net/k900889/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Feb 27, 1990
Decision date	Mar 23, 1990
Days to decision	24 days
Third-party review	No

**APPLICANT**

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Company	<b>Siemens Medical Solutions USA, Inc.</b>
Location	Hoffman Estates, IL, US
Contact	SCOTT A HENSLEY
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/k900889/>; 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 15, 2026