

**K991652 HIGH ENDS GRADIENTS/MAGNETOM SYMPHONY SYSTEM**Aug 3, 1999  
82 days to decisionK991652 · Product code: **LNH** · Radiology  
Source: <https://www.510kdatabase.net/k991652/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	May 13, 1999
Decision date	Aug 3, 1999
Days to decision	82 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Siemens Corp.</b>
Location	Mchenry, IL, US
Contact	KATHLEEN RUTHERFORD
Website	<a href="http://www.siemens.it/">http://www.siemens.it/</a>
510(k) history	66 submissions · 66 cleared · 1978-2010

Siemens Corp. is a global technology company headquartered in McHenry, US. The company develops medical imaging and diagnostic equipment for healthcare providers worldwide. Siemens has received FDA 510(k) clearances from total submissions. The company's regulatory focus centers on Radiology devices, which represent the dominant category of its cleared portfolio. FDA 510(k) clearances span from 1978 to 2010, establishing a significant historical record in medical device regulation. Recent cleared devices include advanced imaging systems such as CT scanners, MR systems, X-r...

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Device record: <https://www.510kdatabase.net/k991652/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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