

**K232482 MAGNETOM Viato.Mobile**Sep 6, 2023  
21 days to decisionK232482 · Product code: **LNH** · Radiology  
Source: <https://www.510kdatabase.net/k232482/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Aug 16, 2023
Decision date	Sep 6, 2023
Days to decision	21 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Siemens Medical Solutions USA, Inc.</b>
Location	Hoffman Estates, IL, US
Contact	Alina Goodman
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/k232482/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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