

**K250443 MAGNETOM Avanto Fit**Jun 16, 2025  
122 days to decisionK250443 · Product code: **LNH** · Radiology  
Source: <https://www.510kdatabase.net/k250443/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Feb 14, 2025
Decision date	Jun 16, 2025
Days to decision	122 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	MAGNETOM Skyra Fit; MAGNETOM Sola Fit; MAGNETOM Viato.Mobile

**APPLICANT**

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Company	<b>Siemens Healthcare GmbH</b>
Location	Erlangen, DE
Contact	Friederike Hertle
510(k) history	30 submissions · 30 cleared · 2016-2026

**REGULATORY CONSULTANT**

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Consulting firm	<b>Siemens Medical Solutions USA, Inc.</b>
Contact	Alina Goodman

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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Device record: <https://www.510kdatabase.net/k250443/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://accessdata.fda.gov)).

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