

K251822 MAGNETOM Free.MaxNov 20, 2025
160 days to decisionK251822 · Product code: **LNH** · Radiology
Source: <https://www.510kdatabase.net/k251822/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Jun 13, 2025
Decision date	Nov 20, 2025
Days to decision	160 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	MAGNETOM Free.Star

APPLICANT

Company	Siemens Shenzhen Magnetic Resonance , Ltd.
Location	Shenzhen, CN
Contact	Li Hai Ting
510(k) history	3 submissions · 3 cleared · 2025-2026

REGULATORY CONSULTANT

Consulting firm	Siemens Medical Solutions USA, Inc.
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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