FDA 510(k) Premarket Notification

K232482 MAGNETOM Viato.Mobile

CLEARED

Sep 6, 2023
21 days to decision

K232482 · Product code: **LNH** · Radiology
Source: https://www.510kdatabase.net/k232482/

SUBMISSION DETAILS

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

Company	**Siemens Medical Solutions USA, Inc.**
Location	Hoffman Estates, IL, US
Contact	Alina Goodman
510(k) history	778 submissions · 778 cleared · 1980-2026