FDA 510(k) Premarket Notification

CLEARED

K192924 MAGNETOM Vida, MAGNETOM Lumina, MAGNETOM Vida Fit

Mar 11, 2020
147 days to decision

K192924 · Product code: **LNH** · Radiology
Source: https://www.510kdatabase.net/k192924/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Oct 16, 2019
Decision date	Mar 11, 2020
Days to decision	147 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	Andrew Turner
510(k) history	779 submissions · 779 cleared · 1980-2026