FDA 510(k) Premarket Notification

K050200 MAGNETOM TRIO A TIM SYSTEM

CLEARED

Feb 28, 2005
31 days to decision

K050200 · Product code: **LNH** · Radiology
Source: https://www.510kdatabase.net/k050200/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Jan 28, 2005
Decision date	Feb 28, 2005
Days to decision	31 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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