FDA 510(k) Premarket Notification

K932271 MAGNETIC RESON DIAG DEVICE MAGNETOM PROJECT 016

CLEARED

Jan 7, 1994
254 days to decision

K932271 · Product code: **LNH** · Radiology
Source: https://www.510kdatabase.net/k932271/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Apr 28, 1993
Decision date	Jan 7, 1994
Days to decision	254 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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