FDA 510(k) Premarket Notification

K961121 KINEMATIC KNEE DEVICE/MAGNETOM OPEN SYSTEM

CLEARED

Jun 19, 1996
90 days to decision

K961121 · Product code: **LNH** · Radiology
Source: https://www.510kdatabase.net/k961121/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Mar 21, 1996
Decision date	Jun 19, 1996
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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