

K961121 KINEMATIC KNEE DEVICE/MAGNETOM OPEN SYSTEMJun 19, 1996
90 days to decisionK961121 · 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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k961121/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 15, 2026