

K030520 INTERA FAMILYMar 6, 2003
15 days to decisionK030520 · Product code: **LNH** · Radiology
Source: <https://www.510kdatabase.net/k030520/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Feb 19, 2003
Decision date	Mar 6, 2003
Days to decision	15 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	Philips Medical Systems North America Co.
Location	Shelton, CT, US
Contact	LYNN HARMER
510(k) history	24 submissions · 24 cleared · 2001-2010

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k030520/> 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 17, 2026