FDA 510(k) Premarket Notification

K030520 INTERA FAMILY

CLEARED

Mar 6, 2003
15 days to decision

K030520 · 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