

**K945397 ANGIO II**Feb 9, 1995  
98 days to decisionK945397 · Product code: **LNH** · Radiology  
Source: <https://www.510kdatabase.net/k945397/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Nov 3, 1994
Decision date	Feb 9, 1995
Days to decision	98 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Philips Medical Systems (Cleveland), Inc.</b>
Location	Mchenry, IL, US
Contact	ELAINE KEELER
510(k) history	190 submissions · 190 cleared · 1977-2017

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k945397/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 16, 2026