

**K133692 IMRI 1.5T AND IMRI 3T S**Feb 10, 2014  
70 days to decisionK133692 · Product code: **LNH** · Radiology  
Source: <https://www.510kdatabase.net/k133692/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	System, Nuclear Magnetic Resonance Imaging (LNH)
Date received	Dec 2, 2013
Decision date	Feb 10, 2014
Days to decision	70 days
Third-party review	Yes
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Imris, Inc.</b>
Location	Rockville, MD, US
Contact	Sanjay Shah
510(k) history	14 submissions · 14 cleared · 2006-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k133692/> 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