

**K013592 EXTREMITY COIL**Jan 8, 2002  
70 days to decisionK013592 · Product code: **MOS** · Radiology  
Source: <https://www.510kdatabase.net/k013592/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Coil, Magnetic Resonance, Specialty (MOS)
Date received	Oct 30, 2001
Decision date	Jan 8, 2002
Days to decision	70 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Usa Instruments, Inc.</b>
Location	Aurora, OH, US
Contact	RONY THOMAS
510(k) history	64 submissions · 64 cleared · 1997-2005

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k013592/> 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 18, 2026