

**K030042 PREMIER III PHASED ARRAY CTL SPINE COIL**Feb 24, 2003  
49 days to decisionK030042 · Product code: **MOS** · Radiology  
Source: <https://www.510kdatabase.net/k030042/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Coil, Magnetic Resonance, Specialty (MOS)
Date received	Jan 6, 2003
Decision date	Feb 24, 2003
Days to decision	49 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Usa Instruments, Inc.</b>
Location	Aurora, OH, US
Contact	CHRISTIE SHUMAKER
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/k030042/> 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