

**K993719 KAC-300 KNEE ARRAY COIL**Nov 26, 1999  
23 days to decisionK993719 · Product code: **MOS** · Radiology  
Source: <https://www.510kdatabase.net/k993719/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Coil, Magnetic Resonance, Specialty (MOS)
Date received	Nov 3, 1999
Decision date	Nov 26, 1999
Days to decision	23 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Mri Devices Corp.</b>
Location	Hartland, WI, US
Contact	THOMAS SCHUBERT
510(k) history	71 submissions · 71 cleared · 1991-2004

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k993719/> 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