

**K093833 KAPSS SPINAL SYSTEM**Oct 15, 2010  
305 days to decisionK093833 · Product code: **MNI** · Orthopedic  
Source: <https://www.510kdatabase.net/k093833/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthosis, Spinal Pedicle Fixation (MNI)
Date received	Dec 14, 2009
Decision date	Oct 15, 2010
Days to decision	305 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Robert Reid, Inc.</b>
Location	Chesterland, OH, US
Contact	KAREN E WARDEN
510(k) history	2 submissions · 2 cleared · 2009-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k093833/> 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