

**K092420 KRD1 PEDICLE SCREW SYSTEM**Nov 19, 2009  
105 days to decisionK092420 · Product code: **MNI** · Orthopedic  
Source: <https://www.510kdatabase.net/k092420/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthosis, Spinal Pedicle Fixation (MNI)
Date received	Aug 6, 2009
Decision date	Nov 19, 2009
Days to decision	105 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Spinefrontier, Inc.</b>
Location	Beverly, MA, US
Contact	JOHN SULLIVAN
510(k) history	24 submissions · 24 cleared · 2007-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k092420/> 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 June 15, 2026