

**K073531 ACCIN PEDICLE SCREW SYSTEM**Feb 4, 2008  
49 days to decisionK073531 · Product code: **MNH** · Orthopedic  
Source: <https://www.510kdatabase.net/k073531/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthosis, Spondylolisthesis Spinal Fixation (MNH)
Date received	Dec 17, 2007
Decision date	Feb 4, 2008
Days to decision	49 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Accelerated Innovation, LLC</b>
Location	Clifton, NJ, US
Contact	MICHAEL KVITNITSKY
510(k) history	7 submissions · 7 cleared · 2008-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k073531/> 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 19, 2026