

**K072672 SEQUOIA SPINAL SYSTEM, MODEL 3300 SERIES**Nov 28, 2007  
68 days to decisionK072672 · Product code: **NKB** · Orthopedic  
Source: <https://www.510kdatabase.net/k072672/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Sep 21, 2007
Decision date	Nov 28, 2007
Days to decision	68 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Abbott Spine, Inc.</b>
Location	Austin, TX, US
Contact	DAVID PADGETT
510(k) history	13 submissions · 13 cleared · 2005-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k072672/> 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 14, 2026