

**K073562 MODIFICATION TO VIPER SPINE SYSTEM**May 15, 2008  
148 days to decisionK073562 · Product code: **NKB** · Orthopedic  
Source: <https://www.510kdatabase.net/k073562/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Dec 19, 2007
Decision date	May 15, 2008
Days to decision	148 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Depuy Spine, Inc.</b>
Location	Raynham, MA, US
Contact	CHRISTOPHER KLACZYK
510(k) history	68 submissions · 67 cleared · 2004-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k073562/> 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 19, 2026