

**K151974 Polaris Spinal System**Nov 13, 2015  
120 days to decisionK151974 · Product code: **NKB** · Orthopedic  
Source: <https://www.510kdatabase.net/k151974/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Jul 16, 2015
Decision date	Nov 13, 2015
Days to decision	120 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Biomet Spine</b>
Location	Warsaw, IN, US
Contact	ALEXANDRA BECK
510(k) history	19 submissions · 18 cleared · 2007-2016

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k151974/> 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 17, 2026