

**K161914 Genesys Spine TiLock Cortical Spinal System**Aug 9, 2016  
28 days to decisionK161914 · Product code: **NKB** · Orthopedic  
Source: <https://www.510kdatabase.net/k161914/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Jul 12, 2016
Decision date	Aug 9, 2016
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Genesys Spine</b>
Location	Austin, TX, US
Contact	William W. Sowers
510(k) history	31 submissions · 31 cleared · 2010-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k161914/> 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