

**K160820 PressON Pro Spinal Fixation System**Jun 28, 2016  
96 days to decisionK160820 · Product code: **NKB** · Orthopedic  
Source: <https://www.510kdatabase.net/k160820/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Mar 24, 2016
Decision date	Jun 28, 2016
Days to decision	96 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Nexus Spine, LLC</b>
Location	Salt Lake City, UT, US
Contact	Jared Crocker
Website	<a href="https://nexusspine.com">https://nexusspine.com</a>
510(k) history	17 submissions · 17 cleared · 2014-2025

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k160820/> 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 26, 2026