

**K183550 Vitality® Spinal Fixation System**Apr 8, 2019  
109 days to decisionK183550 · Product code: **NKB** · Orthopedic  
Source: <https://www.510kdatabase.net/k183550/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Dec 20, 2018
Decision date	Apr 8, 2019
Days to decision	109 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Zimmer Biomet Spine, Inc.</b>
Location	Broomfield, CO, US
Contact	Brendon Gareth Heyerdahl
510(k) history	15 submissions · 15 cleared · 2017-2021

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k183550/> 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