

**K192396 Streamline MIS Spinal Fixation System**Nov 26, 2019  
84 days to decisionK192396 · Product code: **NKB** · Orthopedic  
Source: <https://www.510kdatabase.net/k192396/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Thoracolumbosacral Pedicle Screw System (NKB)
Date received	Sep 3, 2019
Decision date	Nov 26, 2019
Days to decision	84 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Pioneer Surgical Technology, Inc.</b>
Location	Marquette, MI, US
Contact	Michael Mach
510(k) history	26 submissions · 26 cleared · 2010-2025

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k192396/> 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 14, 2026