

**K120724 PIONEER LATERAL PLATE SYSTEM**May 7, 2012  
59 days to decisionK120724 · Product code: **KWQ** · Orthopedic  
Source: <https://www.510kdatabase.net/k120724/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Mar 9, 2012
Decision date	May 7, 2012
Days to decision	59 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k120724/> 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 18, 2026