

**K131724 PIONEER MAXFUSE VERTEBRAL BODY REPLACEMENT DEVICE**Dec 24, 2013  
195 days to decisionK131724 · Product code: **MQP** · Orthopedic  
Source: <https://www.510kdatabase.net/k131724/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spinal Vertebral Body Replacement Device (MQP)
Date received	Jun 12, 2013
Decision date	Dec 24, 2013
Days to decision	195 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	EMILY M DOWNS
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/k131724/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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