

**K073177 PIONEER INTERVERTEBRAL FUSION DEVICE SYSTEM**May 13, 2008  
182 days to decisionK073177 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k073177/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Nov 13, 2007
Decision date	May 13, 2008
Days to decision	182 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Pioneer Surgical Technology</b>
Location	Marquette, MI, US
Contact	JONATHAN M GILBERT
510(k) history	50 submissions · 48 cleared · 1993-2011

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k073177/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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