

**K080673 INTERFUSE INTERVERTEBRAL BODY FUSION  
DEVICE**Jun 10, 2008  
92 days to decisionK080673 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k080673/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Mar 10, 2008
Decision date	Jun 10, 2008
Days to decision	92 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vertebral Technologies, Inc.</b>
Location	Andover, MN, US
Contact	PHILIP B JARVI
510(k) history	7 submissions · 7 cleared · 2008-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k080673/>; 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