

**K093675 INTERFUSE INVERTEBRAL BODY FUSION DEVICE  
MODEL 9076**Dec 23, 2009  
26 days to decisionK093675 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k093675/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Nov 27, 2009
Decision date	Dec 23, 2009
Days to decision	26 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vertebral Technologies, Inc.</b>
Location	Andover, MN, US
Contact	SURESH GHAI
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/k093675/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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