

**K110226 INTERFUSE T INVERTEBRAL BODY FUSION DEVICE**May 27, 2011  
122 days to decisionK110226 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k110226/>**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	Jan 25, 2011
Decision date	May 27, 2011
Days to decision	122 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/k110226/> 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 17, 2026