

**K072289 SPINEFRONTIER DORADO VBR**Nov 1, 2007  
77 days to decisionK072289 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k072289/>**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	Aug 16, 2007
Decision date	Nov 1, 2007
Days to decision	77 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Spinefrontier, Inc.</b>
Location	Beverly, MA, US
Contact	TOM CARLSON
510(k) history	24 submissions · 24 cleared · 2007-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k072289/> 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 June 15, 2026