

**K142347 Amedica Valeo II Interbody Fusion Device**Nov 18, 2014  
88 days to decisionK142347 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k142347/>**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 22, 2014
Decision date	Nov 18, 2014
Days to decision	88 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Amedica Corp.</b>
Location	Salt Lake City, UT, US
Contact	William D Jordan
510(k) history	16 submissions · 16 cleared · 2002-2018

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k142347/> 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 16, 2026