

K143518 Valeo Spacer System and Valeo II Interbody Fusion Device SystemOct 2, 2015
294 days to decisionK143518 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k143518/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Dec 12, 2014
Decision date	Oct 2, 2015
Days to decision	294 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k143518/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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