

**K133583 OPTICAGE INTERBODY FUSION DEVICE, MODEL
SERIES 9070**Feb 10, 2014
81 days to decisionK133583 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k133583/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Nov 21, 2013
Decision date	Feb 10, 2014
Days to decision	81 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Interventional Spine, Inc.
Location	Irvine, CA, US
Contact	JANE METCALF
510(k) history	8 submissions · 7 cleared · 2008-2016

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k133583/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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