

**K091988 MODIFICATION TO: INTERFUSE INTERVERTEBRAL
BODY FUSION DEVICE , MODEL 9076**Jul 30, 2009
28 days to decisionK091988 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k091988/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Jul 2, 2009
Decision date	Jul 30, 2009
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Vertebral Technologies, Inc.
Location	Andover, MN, US
Contact	SURESH GHAI
510(k) history	7 submissions · 7 cleared · 2008-2013

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k091988/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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