

**K101051 AVS A-LAT PEEK SPACERS, MODELS 48750XXX,
48751XXX, 48752XXX, 48753XXX**Aug 12, 2010
119 days to decisionK101051 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k101051/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Apr 15, 2010
Decision date	Aug 12, 2010
Days to decision	119 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Stryker Spine
Location	Allendale, NJ, US
Contact	KIMBERLY S LANE
510(k) history	74 submissions · 73 cleared · 2004-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k101051/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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