

**K073470 AVS PL PEEK SPACERS, MODELS 48351XXX AND
48353XXX**Mar 6, 2008
87 days to decisionK073470 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k073470/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Dec 10, 2007
Decision date	Mar 6, 2008
Days to decision	87 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k073470/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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