

**K091689 SYNAPSE SYSTEM 4.0MM, OC FUSION SYSTEM
4.0MM**

Jul 10, 2009
30 days to decision

K091689 · Product code: **KWP** · Orthopedic
Source: <https://www.510kdatabase.net/k091689/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Interlaminar (KWP)
Date received	Jun 10, 2009
Decision date	Jul 10, 2009
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Synthes (Usa)
Location	Mchenry, IL, US
Contact	STACEY BONNELL
510(k) history	411 submissions · 394 cleared · 1977-2015

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k091689/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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