

K090425 GUIDED LATERAL INTERBODY FUSION (GLIF) SYSTEMMay 19, 2009
89 days to decisionK090425 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k090425/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Feb 19, 2009
Decision date	May 19, 2009
Days to decision	89 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Alphatec Spine, Inc.
Location	Carlsbad, CA, US
Contact	MARY STANNERS
Website	https://www.alphatecspine.com
510(k) history	93 submissions · 93 cleared · 2005-2026

Alphatec Spine, Inc. is a spine surgery medical device company based in Carlsbad, California. The company develops and markets surgical solutions for spinal fusion and fixation procedures. Alphatec Spine maintains a strong FDA 510(k) regulatory record with FDA 510(k) clearances from total submissions. The company specializes in Orthopedic devices, which represent 91% of its submission portfolio. Clearances span from 2005 to 2026, demonstrating sustained regulatory activity and recent market engagement. Recent cleared devices include robotic navigation systems, interbody s...

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Device record: <https://www.510kdatabase.net/k090425/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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