

K210590 Axis Spine Technologies ALIFSep 29, 2021
212 days to decisionK210590 · Product code: **OVD** · Orthopedic
Source: <https://www.510kdatabase.net/k210590/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Integrated Fixation, Lumbar (OVD)
Date received	Mar 1, 2021
Decision date	Sep 29, 2021
Days to decision	212 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Axis Spine Technologies, Ltd.
Location	St. Albans, GB
Contact	Edwin Lindsay
510(k) history	2 submissions · 2 cleared · 2020-2021

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k210590/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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