

K240889 ARTFX Lumbar PEEK CagesOct 2, 2024
184 days to decisionK240889 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k240889/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Apr 1, 2024
Decision date	Oct 2, 2024
Days to decision	184 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Artfx Medical
Location	Jacksonville, FL, US
Contact	Ozgen Ozfidan
510(k) history	3 submissions · 3 cleared · 2024-2024

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k240889/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated May 14, 2026