

K231140 aprevo® transforaminal lumbar interbody fusion deviceMay 19, 2023
28 days to decisionK231140 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k231140/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Apr 21, 2023
Decision date	May 19, 2023
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Carlsmed, Inc.
Location	La Jolla, CA, US
Contact	Karen Liu
510(k) history	20 submissions · 20 cleared · 2020-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k231140/>. 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 13, 2026