

**K232282 aprevo® anterior and lateral lumbar interbody fusion device, aprevo® transforaminal lumbar interbody fusion device**Mar 22, 2024  
235 days to decisionK232282 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k232282/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision              | Substantially Equivalent (Cleared)                         |
| Submission type       | Traditional  |
| Device classification | Intervertebral Fusion Device With Bone Graft, Lumbar (MAX) |
| Date received         | Jul 31, 2023   |
| Decision date         | Mar 22, 2024   |
| Days to decision      | 235 days   |
| Third-party review    | No   |
| Combination product   | No   |
| PCCP authorized       | No   |
| Summary / Statement   | Summary  |

**APPLICANT**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k232282/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 14, 2026