

K171724 Rampart L Lumbar Interbody Fusion DeviceJul 24, 2017
42 days to decisionK171724 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k171724/>**SUBMISSION DETAILS**

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|-----------------------|--|
| Decision | Substantially Equivalent (Cleared) |
| Submission type | Special |
| Device classification | Intervertebral Fusion Device With Bone Graft, Lumbar (MAX) |
| Date received | Jun 12, 2017 |
| Decision date | Jul 24, 2017 |
| Days to decision | 42 days |
| Third-party review | No |
| Summary / Statement | Summary |

APPLICANT

| | |
|----------------|---|
| Company | Spineology, Inc. |
| Location | Stillwater, MN, US |
| Contact | Jacqueline A. Hauge |
| 510(k) history | 54 submissions · 51 cleared · 1999-2025 |

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k171724/> 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 16, 2026