

**K163409 Rampart(TM) T Lumbar Interbody Fusion Device**Mar 8, 2017  
93 days to decisionK163409 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k163409/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Dec 5, 2016
Decision date	Mar 8, 2017
Days to decision	93 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k163409/> 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 16, 2026