

K153082 Rampart A Lumbar Interbody Fusion DeviceFeb 4, 2016
104 days to decisionK153082 · Product code: **MAX** · Orthopedic
Source: <https://www.510kdatabase.net/k153082/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Oct 23, 2015
Decision date	Feb 4, 2016
Days to decision	104 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/k153082/> 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 17, 2026