

**K191367 Harrier-SA Lumbar Interbody System**Jul 5, 2019  
44 days to decisionK191367 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k191367/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	May 22, 2019
Decision date	Jul 5, 2019
Days to decision	44 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Choice Spine, LLC</b>
Location	Knoxville, TN, US
Contact	Kim Finch
510(k) history	9 submissions · 9 cleared · 2019-2023

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k191367/> 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 26, 2026