

**K173832 InTice™-C Porous Ti Cervical Interbody System**May 17, 2018  
150 days to decisionK173832 · Product code: **ODP** · Orthopedic  
Source: <https://www.510kdatabase.net/k173832/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Cervical (ODP)
Date received	Dec 18, 2017
Decision date	May 17, 2018
Days to decision	150 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>X-Spine Systems, Inc.</b>
Location	Centerville, OH, US
Contact	Charlene Brumbaugh
510(k) history	34 submissions · 34 cleared · 2005-2018

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k173832/> 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