

**K173535 Endoskeleton® TAS Interbody Fusion Device,
Endoskeleton® TAS Hyperlordotic Interbody Fusion Device and
Endoskeleton® TCS Interbody Fusion Device**

Feb 13, 2018
90 days to decision

K173535 · Product code: **ODP** · Orthopedic
Source: <https://www.510kdatabase.net/k173535/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Cervical (ODP)
Date received	Nov 15, 2017
Decision date	Feb 13, 2018
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Titan Spine, LLC
Location	Mequon, WI, US
Contact	Jane Rodd
510(k) history	14 submissions · 14 cleared · 2008-2018

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k173535/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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