

# K163494 Summit Spine Channel Cervical Interbody Fusion System

Jun 20, 2017  
189 days to decision

K163494 · Product code: **ODP** · Orthopedic  
Source: <https://www.510kdatabase.net/k163494/>

## SUBMISSION DETAILS

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Cervical (ODP)
Date received	Dec 13, 2016
Decision date	Jun 20, 2017
Days to decision	189 days
Third-party review	No
Summary / Statement	Summary

## APPLICANT

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Company	<b>Summit Spine</b>
Location	Georgetown, TX, US
Contact	Eric Buescher
510(k) history	2 submissions · 2 cleared · 2017-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k163494/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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