

**K100820 WENZEL SPINE VARILIFT INTERBODY FUSION SYSTEM**Aug 5, 2010  
135 days to decisionK100820 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k100820/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Mar 23, 2010
Decision date	Aug 5, 2010
Days to decision	135 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Wenzel Spine</b>
Location	Austin, TX, US
Contact	SOURABH MISHRA
510(k) history	3 submissions · 3 cleared · 2010-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k100820/>. Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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