

**K152651 AccuLIF TL and PL Cage**Dec 15, 2015  
90 days to decisionK152651 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k152651/>**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	Sep 16, 2015
Decision date	Dec 15, 2015
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Stryker Corporation</b>
Location	Malwah, NJ, US
Contact	AAKASH JAIN
Website	<a href="http://www.stryker.com/">http://www.stryker.com/</a>
510(k) history	81 submissions · 81 cleared · 2010-2023

Stryker Corporation is an American multinational medical technology company headquartered in Portage, Michigan. The company develops and markets surgical equipment, neurotechnology, orthopedic implants, and patient safety systems used globally across medical specialties. Stryker has received FDA 510(k) clearances from total submissions between 2010 and 2023. The company's cleared devices span orthopedic surgery, neurosurgery, general and plastic surgery, and ear, nose, and throat specialties. This regulatory record reflects the company's broad portfolio across surgical an...

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