

**K130274 BIOLIGN ACIF SYSTEMS, BIOLIGN STACC ACIF SYSTEM, BIOLIGN TLIF SYSTEM**Oct 29, 2013  
267 days to decisionK130274 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k130274/>**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	Feb 4, 2013
Decision date	Oct 29, 2013
Days to decision	267 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Elite Surgical Supplies (Pty), Ltd.</b>
Location	Marietta, GA, US
Contact	JOHN O'&TOOLE
510(k) history	4 submissions · 4 cleared · 2007-2020

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k130274/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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