

**K131671 MECTALIF**Jul 5, 2013  
28 days to decisionK131671 · Product code: **MAX** · Orthopedic  
Source: <https://www.510kdatabase.net/k131671/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Intervertebral Fusion Device With Bone Graft, Lumbar (MAX)
Date received	Jun 7, 2013
Decision date	Jul 5, 2013
Days to decision	28 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medacta International</b>
Location	San Diego, CA, US
Contact	ADAM GROSS
510(k) history	39 submissions · 39 cleared · 2011-2016

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k131671/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 17, 2026