

**K093629 SOLITAIRE AND SOLITAIRE PEEK ANTERIOR SPINAL SYSTEM**Mar 9, 2010  
106 days to decisionK093629 · Product code: **OVD** · Orthopedic  
Source: <https://www.510kdatabase.net/k093629/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Intervertebral Fusion Device With Integrated Fixation, Lumbar (OVD)
Date received	Nov 23, 2009
Decision date	Mar 9, 2010
Days to decision	106 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Biomet Spine</b>
Location	Warsaw, IN, US
Contact	VIVIAN KELLY
510(k) history	19 submissions · 18 cleared · 2007-2016

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k093629/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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