

**K133802 AIS MODULIFT VBR SYSTEM**Apr 17, 2014  
125 days to decisionK133802 · Product code: **MQP** · Orthopedic  
Source: <https://www.510kdatabase.net/k133802/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spinal Vertebral Body Replacement Device (MQP)
Date received	Dec 13, 2013
Decision date	Apr 17, 2014
Days to decision	125 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Aesculap Implant System, Inc.</b>
Location	Center Valley, PA, US
Contact	LISA M BOYLE
510(k) history	18 submissions · 18 cleared · 2007-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k133802/> 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 14, 2026