

**K060762 AESCULAP PEEK VERTEBRAL BODY REPLACEMENT SYSTEM**Sep 14, 2006  
177 days to decisionK060762 · Product code: **MQP** · Orthopedic  
Source: <https://www.510kdatabase.net/k060762/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spinal Vertebral Body Replacement Device (MQP)
Date received	Mar 21, 2006
Decision date	Sep 14, 2006
Days to decision	177 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Aesculap, Inc.</b>
Location	Burlingame, CA, US
Contact	MATTHEW M HULL
510(k) history	207 submissions · 201 cleared · 1991-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k060762/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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