

**K032059 MODIFICATION TO MACS MODULAR ANTERIOR  
CONSTRUCT SYSTEM**Jul 25, 2003  
23 days to decisionK032059 · Product code: **KWQ** · Orthopedic  
Source: <https://www.510kdatabase.net/k032059/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Jul 2, 2003
Decision date	Jul 25, 2003
Days to decision	23 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Aesculap, Inc.</b>
Location	Burlingame, CA, US
Contact	GEORG KELLER
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/k032059/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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