

**K050804 MODIFICATION TO: SPECTRUM™ CERVICAL SPINAL SYSTEM**Apr 11, 2005  
12 days to decisionK050804 · Product code: **KWQ** · Orthopedic  
Source: <https://www.510kdatabase.net/k050804/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Intervertebral Body (KWQ)
Date received	Mar 30, 2005
Decision date	Apr 11, 2005
Days to decision	12 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Aesculap, Inc.</b>
Location	Burlingame, CA, US
Contact	KATHY A RACOSKY
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/k050804/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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