

**K051697 SELECTIVE-AXIS POSTERIOR NONCERVICAL PLATING SYSTEM**

Sep 21, 2005  
90 days to decision

K051697 · Product code: **MNH** · Orthopedic  
Source: <https://www.510kdatabase.net/k051697/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Orthosis, Spondylolisthesis Spinal Fixation (MNH)
Date received	Jun 23, 2005
Decision date	Sep 21, 2005
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Medical Device Advisory Development Group</b>
Location	Irvine, CA, US
Contact	C. STEPHEN LAWRENCE
510(k) history	1 submissions · 1 cleared · 2005-2005

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k051697/>; Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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