

**K990845 MODIFICATION TO PARTS NUMBERED CGT-SL AND CGT-SL-15**

Apr 13, 1999  
29 days to decision

K990845 · Product code: **KWP** · Orthopedic  
Source: <https://www.510kdatabase.net/k990845/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Appliance, Fixation, Spinal Interlaminar (KWP)
Date received	Mar 15, 1999
Decision date	Apr 13, 1999
Days to decision	29 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Advanced Spine Fixation Systems, Inc.</b>
Location	Washington, DC, US
Contact	GREG HOLLAND
510(k) history	16 submissions · 7 cleared · 1989-2000

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

Device record: <https://www.510kdatabase.net/k990845/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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