

**K050767 STERLING INTERFERENCE SCREW ST**Jun 9, 2005  
76 days to decisionK050767 · Product code: **HWC** · Orthopedic  
Source: <https://www.510kdatabase.net/k050767/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Screw, Fixation, Bone (HWC)
Date received	Mar 25, 2005
Decision date	Jun 9, 2005
Days to decision	76 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Regeneration Technologies, Inc.</b>
Location	Alachua, FL, US
Contact	CARRIE HARTILL
510(k) history	11 submissions · 11 cleared · 2005-2008

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k050767/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 19, 2026