

**K081813 UNIVERSAL COMPRESSION FRACTURE REPAIR SYSTEM**Mar 12, 2009  
259 days to decisionK081813 · Product code: **HWC** · Orthopedic  
Source: <https://www.510kdatabase.net/k081813/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Screw, Fixation, Bone (HWC)
Date received	Jun 26, 2008
Decision date	Mar 12, 2009
Days to decision	259 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>U.S. Implant Solutions, LLC</b>
Location	Oviedo, FL, US
Contact	SCOTT GARRETT
510(k) history	1 submissions · 1 cleared · 2009-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k081813/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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