

**K191817 ActivOrtho Continuous Compression Screw System**Jul 26, 2019  
21 days to decisionK191817 · Product code: **HWC** · Orthopedic  
Source: <https://www.510kdatabase.net/k191817/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Screw, Fixation, Bone (HWC)
Date received	Jul 5, 2019
Decision date	Jul 26, 2019
Days to decision	21 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Activortho, Inc.</b>
Location	Plymouth, MN, US
Contact	Paul Hindrichs
510(k) history	2 submissions · 2 cleared · 2019-2019

**REGULATORY CONSULTANT**

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Consulting firm	<b>DuVal &amp; Associates, P.A.</b>
Contact	Lisa L. Pritchard

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k191817/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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