

**K180050 CurvaFix Intramedullary Rodscrew System**Mar 1, 2019  
417 days to decisionK180050 · Product code: **HWC** · Orthopedic  
Source: <https://www.510kdatabase.net/k180050/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Screw, Fixation, Bone (HWC)
Date received	Jan 8, 2018
Decision date	Mar 1, 2019
Days to decision	417 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Curvafix, Inc.</b>
Location	Bellevue, WA, US
Contact	Steve Dimmer
510(k) history	3 submissions · 3 cleared · 2019-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k180050/> 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 26, 2026