

**K200840 DuoHex CH Cannulated Hammertoe System**Jun 26, 2020  
87 days to decisionK200840 · Product code: **HWC** · Orthopedic  
Source: <https://www.510kdatabase.net/k200840/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Screw, Fixation, Bone (HWC)
Date received	Mar 31, 2020
Decision date	Jun 26, 2020
Days to decision	87 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Nextremity Solutions, Inc.</b>
Location	Red Bank, NJ, US
Contact	Elise Fox
510(k) history	17 submissions · 17 cleared · 2014-2022

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k200840/> 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