

**K200951 Accu-Joint Hemi Implant**Dec 8, 2020  
243 days to decisionK200951 · Product code: **KWD** · Orthopedic  
Source: <https://www.510kdatabase.net/k200951/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Toe, Hemi-, Phalangeal (KWD)
Date received	Apr 9, 2020
Decision date	Dec 8, 2020
Days to decision	243 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Accufix Surgical, Inc.</b>
Location	West Haven, CT, US
Contact	Michael Parisi
510(k) history	2 submissions · 2 cleared · 2020-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k200951/> 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 27, 2026