

**K072922 3S HEMI TOE**Nov 28, 2007  
44 days to decisionK072922 · Product code: **KWD** · Orthopedic  
Source: <https://www.510kdatabase.net/k072922/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Prosthesis, Toe, Hemi-, Phalangeal (KWD)
Date received	Oct 15, 2007
Decision date	Nov 28, 2007
Days to decision	44 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Trilliant Surgical, Ltd.</b>
Location	Round Rock, TX, US
Contact	JD WEBB
510(k) history	14 submissions · 14 cleared · 2007-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k072922/> 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 16, 2026