

**K130334 HOFFMANN LRF (LIMB RECONSTRUCTION FRAME SYSTEM)**May 3, 2013  
81 days to decisionK130334 · Product code: **KTT** · Orthopedic  
Source: <https://www.510kdatabase.net/k130334/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Nail/blade/plate Combination, Multiple Component (KTT)
Date received	Feb 11, 2013
Decision date	May 3, 2013
Days to decision	81 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Stryker Trauma AG</b>
Location	Malwah, NJ, US
Contact	ESTELA CELI
510(k) history	25 submissions · 25 cleared · 2013-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k130334/>; 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