

**K130907 HOFFMANN LRF (LIMB RECONSTRUCTION FRAME)
SYSTEM**Jun 10, 2013
70 days to decisionK130907 · Product code: **KTT** · Orthopedic
Source: <https://www.510kdatabase.net/k130907/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Appliance, Fixation, Nail/blade/plate Combination, Multiple Component (KTT)
Date received	Apr 1, 2013
Decision date	Jun 10, 2013
Days to decision	70 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k130907/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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