

**K121601 SYNTHES 2.7/3.5MM VARIABLE ANGLE LCP ANKLE TRAUMA SYSTEM-ANTEROLATERAL DISTAL TIBIA PLATE**

Jul 6, 2012  
35 days to decision

K121601 · Product code: **HRS** · Orthopedic  
Source: <https://www.510kdatabase.net/k121601/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plate, Fixation, Bone (HRS)
Date received	Jun 1, 2012
Decision date	Jul 6, 2012
Days to decision	35 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Synthes (Usa)</b>
Location	Mchenry, IL, US
Contact	ANGELA F LASSANDRO
510(k) history	411 submissions · 394 cleared · 1977-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k121601/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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