

**K052390 SYNTHES LCP PROXIMAL TIBIA PLATES LINE EXTENSION**

Oct 7, 2005  
37 days to decision

K052390 · Product code: **HRS** · Orthopedic  
Source: <https://www.510kdatabase.net/k052390/>

**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plate, Fixation, Bone (HRS)
Date received	Aug 31, 2005
Decision date	Oct 7, 2005
Days to decision	37 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Synthes (Usa)</b>
Location	Mchenry, IL, US
Contact	Lisa Boyle
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/k052390/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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