

**K082624 SYNTHES (USA) 3.5MM LCP POSTEROMEDIAL
PROXIMAL TIBIA PLATES**Nov 10, 2008
62 days to decisionK082624 · Product code: **HRS** · Orthopedic
Source: <https://www.510kdatabase.net/k082624/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Plate, Fixation, Bone (HRS)
Date received	Sep 9, 2008
Decision date	Nov 10, 2008
Days to decision	62 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Synthes (Usa)
Location	Mchenry, IL, US
Contact	KARL J NITTINGER
510(k) history	411 submissions · 394 cleared · 1977-2015

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k082624/> Data sourced from FDA 510(k) public records (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. - Generated May 14, 2026