

K150394 Tyber Medical Wedge SystemJul 28, 2015
160 days to decisionK150394 · Product code: **PLF** · Orthopedic
Source: <https://www.510kdatabase.net/k150394/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Bone Wedge (PLF)
Date received	Feb 18, 2015
Decision date	Jul 28, 2015
Days to decision	160 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Tyber Medical, LLC
Location	Morristown, NJ, US
Contact	JEFF TYBER
Website	https://www.tybermed.com
510(k) history	28 submissions · 28 cleared · 2013-2026

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k150394/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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