

**K103145 HAMMERLOC TM TIBIAL FIXATION DEVICE**Feb 9, 2011  
107 days to decisionK103145 · Product code: **MBI** · Orthopedic  
Source: <https://www.510kdatabase.net/k103145/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Fastener, Fixation, Nondegradable, Soft Tissue (MBI)
Date received	Oct 25, 2010
Decision date	Feb 9, 2011
Days to decision	107 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Biomet Sports Medicine</b>
Location	Warsaw, IN, US
Contact	ROBERT FRIDDLE
510(k) history	9 submissions · 9 cleared · 2008-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k103145/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://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](http://accessdata.fda.gov). - Generated May 14, 2026