

**K072063 KFX TISSUE FIXATION SYSTEM AND ACCESSORIES**Oct 26, 2007  
91 days to decisionK072063 · Product code: **MBI** · Orthopedic  
Source: <https://www.510kdatabase.net/k072063/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Fastener, Fixation, Nondegradable, Soft Tissue (MBI)
Date received	Jul 27, 2007
Decision date	Oct 26, 2007
Days to decision	91 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Kfx Medical</b>
Location	Carlsbad, CA, US
Contact	M D HEAVEN
510(k) history	7 submissions · 7 cleared · 2006-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k072063/> 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