

**K131182 MULTIFIX S KNOTLESS FIXATION DEVICE**Jul 29, 2013  
94 days to decisionK131182 · Product code: **MBI** · Orthopedic  
Source: <https://www.510kdatabase.net/k131182/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Fastener, Fixation, Nondegradable, Soft Tissue (MBI)
Date received	Apr 26, 2013
Decision date	Jul 29, 2013
Days to decision	94 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Arthocare Corporation</b>
Location	Harrogate, North Yorkshire, GB
Contact	MITCHELL DHORITY
510(k) history	6 submissions · 6 cleared · 2001-2016

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k131182/> 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