

**K093717 C-MOR VISUALIZATION DEVICE**Mar 2, 2010  
90 days to decisionK093717 · Product code: **HRX** · Orthopedic  
Source: <https://www.510kdatabase.net/k093717/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Dec 2, 2009
Decision date	Mar 2, 2010
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Axis Surgical Technologies, Inc.</b>
Location	Mountain View, CA, US
Contact	JAMES FARNWORTH
510(k) history	1 submissions · 1 cleared · 2010-2010

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k093717/> 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 19, 2026