

**K090815 PERCUTANEOUS DISCECTOMY SYSTEM**May 21, 2009  
57 days to decisionK090815 · Product code: **HRX** · Orthopedic  
Source: <https://www.510kdatabase.net/k090815/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Mar 25, 2009
Decision date	May 21, 2009
Days to decision	57 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Laurimed, LLC</b>
Location	Redwood City, CA, US
Contact	SEVRINA CIUCCI
510(k) history	8 submissions · 8 cleared · 2008-2017

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k090815/> 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