

**K090303 XTEND-ST NUCLEUS REMOVAL SYSTEM, MODEL 9004**Apr 30, 2009  
83 days to decisionK090303 · Product code: **HRX** · Orthopedic  
Source: <https://www.510kdatabase.net/k090303/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Feb 6, 2009
Decision date	Apr 30, 2009
Days to decision	83 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Corespine Technologies, LLC</b>
Location	Minneapolis, MN, US
Contact	BRITT K NORTON
510(k) history	1 submissions · 1 cleared · 2009-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k090303/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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