

**K922391 ENDOSCOPIC CARPAL TUNNEL INSTRUMENTS**Aug 18, 1992  
90 days to decisionK922391 · Product code: **HRX** · Orthopedic  
Source: <https://www.510kdatabase.net/k922391/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	May 20, 1992
Decision date	Aug 18, 1992
Days to decision	90 days
Third-party review	No
Summary / Statement	Statement

**APPLICANT**

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Company	<b>Instratek, Inc.</b>
Location	Houston, TX, US
Contact	PERRY FORRESTER
510(k) history	10 submissions · 10 cleared · 1992-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k922391/> 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 17, 2026