

**K191594 Nuvis Battery Arthroscope**Jul 11, 2019  
24 days to decisionK191594 · Product code: **HRX** · General & Plastic Surgery  
Source: <https://www.510kdatabase.net/k191594/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Arthroscope (HRX)
Date received	Jun 17, 2019
Decision date	Jul 11, 2019
Days to decision	24 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Integrated Endoscopy, Inc.</b>
Location	Rancho Santa Margarita, CA, US
Contact	Albert Rego
510(k) history	3 submissions · 3 cleared · 2014-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k191594/> 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 26, 2026