

**K230205 Veloxion System**Feb 24, 2023  
30 days to decisionK230205 · Product code: **FJL** · Gastroenterology & Urology  
Source: <https://www.510kdatabase.net/k230205/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Special
Device classification	Resectoscope (FJL)
Date received	Jan 25, 2023
Decision date	Feb 24, 2023
Days to decision	30 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Corinth Medtech, Inc.</b>
Location	Cupertino, CA, US
Contact	George Hsu
510(k) history	7 submissions · 7 cleared · 2017-2023

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k230205/> 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 27, 2026