

**K243020 MIDASVu**Dec 2, 2024  
66 days to decisionK243020 · Product code: **HRX** · Orthopedic  
Source: <https://www.510kdatabase.net/k243020/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Arthroscope (HRX)
Date received	Sep 27, 2024
Decision date	Dec 2, 2024
Days to decision	66 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Intravu, Inc.</b>
Location	Redwood City, CA, US
Contact	Jenna Rapoport
510(k) history	2 submissions · 2 cleared · 2018-2024

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k243020/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated May 13, 2026