

**K220567 Easi-Vue embolic microspheres System**Sep 7, 2022  
191 days to decisionK220567 · Product code: **KRD** · CardiovascularSource: <https://www.510kdatabase.net/k220567/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Vascular, For Promoting Embolization (KRD)
Date received	Feb 28, 2022
Decision date	Sep 7, 2022
Days to decision	191 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Abk Biomedical, Inc.</b>
Location	Halifax, CA
Contact	Brandi Woods
510(k) history	2 submissions · 2 cleared · 2022-2024

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k220567/> 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