

**K133924 ENDOLUMINAL OCCLUSION SYSTEM (EOS)**Dec 3, 2014  
345 days to decisionK133924 · Product code: **KRD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k133924/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Abbreviated
Device classification	Device, Vascular, For Promoting Embolization (KRD)
Date received	Dec 23, 2013
Decision date	Dec 3, 2014
Days to decision	345 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

Company	<b>Artventive Medical Group, Inc.</b>
Location	Woodinville, WA, US
Contact	Roberta Hines
510(k) history	3 submissions · 3 cleared · 2014-2022

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k133924/> 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