

**K212057 Endoluminal Occlusion System, EOS-X, Endoluminal Occlusion System, EOS**Mar 1, 2022  
243 days to decisionK212057 · Product code: **KRD** · Cardiovascular  
Source: <https://www.510kdatabase.net/k212057/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Device, Vascular, For Promoting Embolization (KRD)
Date received	Jul 1, 2021
Decision date	Mar 1, 2022
Days to decision	243 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k212057/>. 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