

**K260051 CorVision® (91-205-Z0014)**May 12, 2026  
124 days to decisionK260051 · Product code: **MOD** · AnesthesiologySource: <https://www.510kdatabase.net/k260051/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Accessory To Continuous Ventilator (respirator) (MOD)
Date received	Jan 8, 2026
Decision date	May 12, 2026
Days to decision	124 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Corvent Medical, Inc.</b>
Location	Fargo, ND, US
Contact	Richard Walsh
510(k) history	2 submissions · 2 cleared · 2024-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k260051/> 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 July 6, 2026