

**K251245 OxyMinder Pro (10310)**Aug 11, 2025  
111 days to decisionK251245 · Product code: **CCL** · Anesthesiology  
Source: <https://www.510kdatabase.net/k251245/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Gas, Oxygen, Gaseous-phase (CCL)
Date received	Apr 22, 2025
Decision date	Aug 11, 2025
Days to decision	111 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Bio-Med Device, Inc.</b>
Location	Guilford, CT, US
Contact	Max Le
510(k) history	1 submissions · 1 cleared · 2025-2025

**REGULATORY CONSULTANT**

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Consulting firm	<b>ProMedic Consulting, LLC</b>
Contact	Paul Dryden

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: FDA accessdata.fda.gov

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k251245/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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