

**K254104 ResScan Essentials**May 27, 2026  
159 days to decisionK254104 · Product code: **BZD** · Anesthesiology  
Source: <https://www.510kdatabase.net/k254104/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Non-continuous (respirator) (BZD)
Date received	Dec 19, 2025
Decision date	May 27, 2026
Days to decision	159 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>ResMed Corp</b>
Location	Poway, CA, US
Contact	Marzia Signoroni
Website	<a href="http://www.resmed.com/">http://www.resmed.com/</a>
510(k) history	17 submissions · 17 cleared · 1997-2026

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k254104/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](https://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](https://accessdata.fda.gov). - Generated June 19, 2026