

**K070263 APNEALINK**Jun 15, 2007  
140 days to decisionK070263 · Product code: **MNR** · Anesthesiology  
Source: <https://www.510kdatabase.net/k070263/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilatory Effort Recorder (MNR)
Date received	Jan 26, 2007
Decision date	Jun 15, 2007
Days to decision	140 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Resmed Germany, Inc.</b>
Location	Poway, CA, US
Contact	DAVID D&apos;CRUZ
510(k) history	8 submissions · 8 cleared · 2006-2015

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k070263/> 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 24, 2026