

**K090398 PROVENT PROFESSIONAL SLEEP APNEA THERAPY**Apr 3, 2009  
45 days to decisionK090398 · Product code: **OHP** · Dental  
Source: <https://www.510kdatabase.net/k090398/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Expiratory Resistance Valve, Intranasal, For Obstructive Sleep Apnea (OHP)
Date received	Feb 17, 2009
Decision date	Apr 3, 2009
Days to decision	45 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Ventus Medical, Inc.</b>
Location	Hillsboroguh, CA, US
Contact	MARY ROSE
510(k) history	5 submissions · 5 cleared · 2008-2012

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k090398/> 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