

**K102404 PROVENT PROFESSIONAL SLEEP APNEA THERAPY  
(PROVENT 80)**Dec 2, 2010  
100 days to decisionK102404 · Product code: **OHP** · Dental  
Source: <https://www.510kdatabase.net/k102404/>**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	Aug 24, 2010
Decision date	Dec 2, 2010
Days to decision	100 days
Third-party review	No
Summary / Statement	Summary
Other names	PROVENT PROFESSIONAL SLEEP APNEA THERAPY (PROVENT 50)

**APPLICANT**

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Company	<b>Ventus Medical, Inc.</b>
Location	Hillsboroguh, CA, US
Contact	MICHAEL NEVARES
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/k102404/> 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