

**K992384 CPAP/PRO CPAP INTERFACE**Dec 1, 1999  
138 days to decisionK992384 · Product code: **BZD** · Anesthesiology  
Source: <https://www.510kdatabase.net/k992384/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Non-continuous (respirator) (BZD)
Date received	Jul 16, 1999
Decision date	Dec 1, 1999
Days to decision	138 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Stevenson Industries, Inc.</b>
Location	Simi Valley, CA, US
Contact	DAVID W SCHLERF
510(k) history	2 submissions · 2 cleared · 1999-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k992384/> 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