

**K952880 MODEL LP-1000 VOLUME VENTILATOR**Jun 20, 1996  
364 days to decisionK952880 · Product code: **CBK** · AnesthesiologySource: <https://www.510kdatabase.net/k952880/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent - ST
Submission type	Traditional
Device classification	Ventilator, Continuous, Facility Use (CBK)
Date received	Jun 22, 1995
Decision date	Jun 20, 1996
Days to decision	364 days
Third-party review	No

**APPLICANT**

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Company	<b>Aequitron Medical, Inc.</b>
Location	Walker, MI, US
Contact	ROBERT C SAMEC
510(k) history	24 submissions · 22 cleared · 1984-1997

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k952880/>, 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 25, 2026