

**K900115 VENTLAB EXHALATION MANIFOLD VX-2000 A & VX-2000 B**Feb 22, 1990  
44 days to decisionK900115 · Product code: **CAN** · Anesthesiology  
Source: <https://www.510kdatabase.net/k900115/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Regulator, Pressure, Gas Cylinder (CAN)
Date received	Jan 9, 1990
Decision date	Feb 22, 1990
Days to decision	44 days
Third-party review	No

**APPLICANT**

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Company	<b>Ventlab Corp.</b>
Location	Hackensack, NJ, US
Contact	GREGORY LAU
510(k) history	22 submissions · 22 cleared · 1989-2012

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**510k Database** - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k900115/> 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