

**K882328 HI FLOW 50**Jun 28, 1988  
22 days to decisionK882328 · Product code: **BYJ** · Anesthesiology  
Source: <https://www.510kdatabase.net/k882328/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Liquid-oxygen, Portable (BYJ)
Date received	Jun 6, 1988
Decision date	Jun 28, 1988
Days to decision	22 days
Third-party review	No

**APPLICANT**

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Company	<b>Minnesota Valley Engineering, Inc.</b>
Location	New Prague, MN, US
Contact	SCHOENBAUER
510(k) history	4 submissions · 4 cleared · 1987-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k882328/>; 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 27, 2026