

**K875352 HI FLOW PORTABLE LIQUID OXYGEN UNIT**Feb 29, 1988  
60 days to decisionK875352 · Product code: **BYJ** · AnesthesiologySource: <https://www.510kdatabase.net/k875352/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Liquid-oxygen, Portable (BYJ)
Date received	Dec 31, 1987
Decision date	Feb 29, 1988
Days to decision	60 days
Third-party review	No

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k875352/>; 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