

**K863978 PORTABLE LIQUID OXYGEN UNIT (MODIFICATION)**Dec 15, 1986  
62 days to decisionK863978 · Product code: **BYJ** · Anesthesiology  
Source: <https://www.510kdatabase.net/k863978/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Liquid-oxygen, Portable (BYJ)
Date received	Oct 14, 1986
Decision date	Dec 15, 1986
Days to decision	62 days
Third-party review	No

**APPLICANT**

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Company	<b>Penox Technologies, Inc.</b>
Location	Pittston, PA, US
Contact	THANA A FRANCE
510(k) history	10 submissions · 10 cleared · 1985-1987

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k863978/>; 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