

**K895082 VOYAGER PX**Jan 9, 1990  
148 days to decisionK895082 · Product code: **BYJ** · Anesthesiology  
Source: <https://www.510kdatabase.net/k895082/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Liquid-oxygen, Portable (BYJ)
Date received	Aug 14, 1989
Decision date	Jan 9, 1990
Days to decision	148 days
Third-party review	No

**APPLICANT**

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Company	<b>Smith-Perry Corp.</b>
Location	Riviera Beach, FL, US
Contact	DAVID SMITH
510(k) history	2 submissions · 2 cleared · 1989-1990

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k895082/>; 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 26, 2026