

**K892639 THE ORIGINAL PULSE-PAK**Oct 5, 1989  
176 days to decisionK892639 · Product code: **NFB** · Anesthesiology  
Source: <https://www.510kdatabase.net/k892639/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Conserver, Oxygen (NFB)
Date received	Apr 12, 1989
Decision date	Oct 5, 1989
Days to decision	176 days
Third-party review	No

**APPLICANT**

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Company	<b>Jones Sales Co.</b>
Location	Canal Winchester, OH, US
Contact	RUSSELL, SR.
510(k) history	1 submissions · 1 cleared · 1989-1989

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k892639/>; 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