

**K131990 OXYGEN GENERATOR LIQUEFIER - (OGL)**Feb 21, 2014  
238 days to decisionK131990 · Product code: **CAW** · Anesthesiology  
Source: <https://www.510kdatabase.net/k131990/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Generator, Oxygen, Portable (CAW)
Date received	Jun 28, 2013
Decision date	Feb 21, 2014
Days to decision	238 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Essex Cryogenics of Missouri, Inc.</b>
Location	St. Louis, MO, US
Contact	PAUL DRYDEN
510(k) history	5 submissions · 5 cleared · 1988-2014

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k131990/> 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 24, 2026