

**K013931 OMNI OXYGEN SYSTEM, MODEL 1000**Mar 27, 2002  
119 days to decisionK013931 · Product code: **CAW** · Anesthesiology  
Source: <https://www.510kdatabase.net/k013931/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Generator, Oxygen, Portable (CAW)
Date received	Nov 28, 2001
Decision date	Mar 27, 2002
Days to decision	119 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Sequal Technologies, Inc.</b>
Location	San Diego, CA, US
Contact	PAM JACKSON
510(k) history	4 submissions · 4 cleared · 2001-2009

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k013931/> 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