

**K001708 PROFOX SOFTWARE, MODEL 920M-41**Mar 1, 2001  
269 days to decisionK001708 · Product code: **DQA** · Anesthesiology  
Source: <https://www.510kdatabase.net/k001708/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Jun 5, 2000
Decision date	Mar 1, 2001
Days to decision	269 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Respironics Georgia, Inc.</b>
Location	Marietta, GA, US
Contact	GERRY RICHARDSON
510(k) history	2 submissions · 2 cleared · 2001-2002

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k001708/> 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 25, 2026