

**K874198 VITALOG HMS-3000**Mar 22, 1988  
165 days to decisionK874198 · Product code: **DQA** · AnesthesiologySource: <https://www.510kdatabase.net/k874198/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Oct 9, 1987
Decision date	Mar 22, 1988
Days to decision	165 days
Third-party review	No

**APPLICANT**

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Company	<b>Vitalog Corp.</b>
Location	Walker, MI, US
Contact	BRUCE RULE
510(k) history	4 submissions · 4 cleared · 1981-1988

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k874198/>; 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