

**K012344 VANGUARD REPROCESSED PULSE OXIMETER  
SENSORS**Oct 4, 2002  
437 days to decisionK012344 · Product code: **NLF** · Anesthesiology  
Source: <https://www.510kdatabase.net/k012344/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter, Reprocessed (NLF)
Date received	Jul 24, 2001
Decision date	Oct 4, 2002
Days to decision	437 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Vanguard Medical Concepts, Inc.</b>
Location	Plant City, FL, US
Contact	MIKE SAMMON
510(k) history	33 submissions · 33 cleared · 1991-2005

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k012344/> 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 20, 2026