

**K051352 OXIMAX NPB-40 HANDHELD PULSE OXIMETER**Aug 11, 2005  
79 days to decisionK051352 · Product code: **DQA** · AnesthesiologySource: <https://www.510kdatabase.net/k051352/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	May 24, 2005
Decision date	Aug 11, 2005
Days to decision	79 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Nellcor Puritan Bennett, Inc.</b>
Location	Minneapolis, MN, US
Contact	SARAH HARRINGTON
510(k) history	42 submissions · 37 cleared · 1996-2007

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k051352/> 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