

**K123581 NELLCOR OXIMAX N-600X PULSE OXIMETER,  
NELLCOR BEDSIDE RESPIRATORY PATIENT MONITORING  
SYSTEM, NELLCOR BEDSIDE SPO2 PATIEN**May 9, 2013  
170 days to decisionK123581 · Product code: **DQA** · Anesthesiology  
Source: <https://www.510kdatabase.net/k123581/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Nov 20, 2012
Decision date	May 9, 2013
Days to decision	170 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Covidien, Formerly Nellcor Puritan Bennett, Inc.</b>
Location	Boulder, CO, US
Contact	MIA M WARE
510(k) history	5 submissions · 5 cleared · 2008-2013

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k123581/> 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