

**K012891 OXIMAX PULSE OXIMETRY SYSTEM WITH N-595  
PULSE OXIMETER AND OXIMAX SENSORS AND CABLES (AKA  
ACCESSORIES)**Mar 7, 2002  
191 days to decisionK012891 · Product code: **DQA** · Anesthesiology  
Source: <https://www.510kdatabase.net/k012891/>**SUBMISSION DETAILS**

---

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Aug 28, 2001
Decision date	Mar 7, 2002
Days to decision	191 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

---

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

---

510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k012891/> 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