

**K033973 OXISENSOR II RECYCLED SENSOR AND OXIMAX
RECYCLED SENSOR**Dec 13, 2004
357 days to decisionK033973 · Product code: **NLF** · Anesthesiology
Source: <https://www.510kdatabase.net/k033973/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter, Reprocessed (NLF)
Date received	Dec 22, 2003
Decision date	Dec 13, 2004
Days to decision	357 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Nellcor Puritan Bennett, Inc.
Location	Minneapolis, MN, US
Contact	LUANNE NG
510(k) history	42 submissions · 37 cleared · 1996-2007

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k033973/> Data sourced from FDA 510(k) public records (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. - Generated May 25, 2026