

**K012609 CLEARMEDICAL/NELLCOR OXISENSOR II,
NEONATAL/ADULT, MODEL N-25**Jul 3, 2002
324 days to decisionK012609 · Product code: **NLF** · Anesthesiology
Source: <https://www.510kdatabase.net/k012609/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter, Reprocessed (NLF)
Date received	Aug 13, 2001
Decision date	Jul 3, 2002
Days to decision	324 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Clearmedical, Inc.
Location	Bellevue, WA, US
Contact	RICHARD RADFORD
510(k) history	14 submissions · 14 cleared · 2002-2005

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k012609/> 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