FDA 510(k) Premarket Notification

K012609 CLEARMEDICAL/NELLCOR OXISENSOR II, NEONATAL/ADULT, MODEL N-25

CLEARED

Jul 3, 2002
324 days to decision

K012609 · 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