FDA 510(k) Premarket Notification

K043359 4100 PATIENT OXIMETER MODULE

CLEARED

Jan 7, 2005
32 days to decision

K043359 · Product code: **DQA** · Anesthesiology
Source: https://www.510kdatabase.net/k043359/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Dec 6, 2004
Decision date	Jan 7, 2005
Days to decision	32 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	**Nonin Medical, Inc.**
Location	White Bear Lake, MN, US
Contact	JOHN R DALPEE
510(k) history	50 submissions · 50 cleared · 1986-2024

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k043359/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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