FDA 510(k) Premarket Notification

K890033 MODEL 8700 PULSE OXIMETER

CLEARED

Feb 6, 1989
33 days to decision

K890033 · Product code: **DQA** · Anesthesiology
Source: https://www.510kdatabase.net/k890033/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Jan 4, 1989
Decision date	Feb 6, 1989
Days to decision	33 days
Third-party review	No

APPLICANT

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

510k Database - www.510kdatabase.net
Device record: https://www.510kdatabase.net/k890033/ Data sourced from FDA 510(k) public records (accessdata.fda.gov).
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