FDA 510(k) Premarket Notification

K051352 OXIMAX NPB-40 HANDHELD PULSE OXIMETER

CLEARED

Aug 11, 2005
79 days to decision

K051352 · Product code: **DQA** · Anesthesiology
Source: https://www.510kdatabase.net/k051352/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	May 24, 2005
Decision date	Aug 11, 2005
Days to decision	79 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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