FDA 510(k) Premarket Notification

K033973 OXISENSOR II RECYCLED SENSOR AND OXIMAX RECYCLED SENSOR

CLEARED

Dec 13, 2004
357 days to decision

K033973 · Product code: **NLF** · Anesthesiology
Source: https://www.510kdatabase.net/k033973/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter, Reprocessed (NLF)
Date received	Dec 22, 2003
Decision date	Dec 13, 2004
Days to decision	357 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

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