

K092368 REPROCESSED PULSE OXIMETER SENSORSNov 3, 2009
90 days to decisionK092368 · Product code: **NLF** · Anesthesiology
Source: <https://www.510kdatabase.net/k092368/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter, Reprocessed (NLF)
Date received	Aug 5, 2009
Decision date	Nov 3, 2009
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Sterilmed, Inc.
Location	Plymouth, MN, US
Contact	GARRETT AHLBORG
510(k) history	64 submissions · 64 cleared · 2001-2024

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