

**K241758 Reprocessed RD SET Adt Pulse Oximeter Sensor (4000)**Oct 22, 2024  
124 days to decisionK241758 · Product code: **NLF** · Anesthesiology  
Source: <https://www.510kdatabase.net/k241758/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter, Reprocessed (NLF)
Date received	Jun 20, 2024
Decision date	Oct 22, 2024
Days to decision	124 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary
Other names	Reprocessed RD SET Pdt Pulse Oximeter Sensor (4001); Reprocessed RD SET Inf Pulse Oximeter Sensor (4002); Reprocessed RD SET Adt Pulse Oximeter Sensor (4003)

**APPLICANT**

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Company	<b>Stryker Sustainability Solutions</b>
Location	Tempe, AZ, US
Contact	Mia Brown
510(k) history	31 submissions · 31 cleared · 2011-2025

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k241758/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at [accessdata.fda.gov](http://accessdata.fda.gov). - Generated May 13, 2026