

**K111773 RENU MEDICAL REPROCESSED OXIMAX SENSORS:  
ADULT, PEDIATRIC, INFANT AND NEONATE**Nov 16, 2011  
146 days to decisionK111773 · Product code: **NLF** · Anesthesiology  
Source: <https://www.510kdatabase.net/k111773/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter, Reprocessed (NLF)
Date received	Jun 23, 2011
Decision date	Nov 16, 2011
Days to decision	146 days
Third-party review	No
Summary / Statement	Summary

**APPLICANT**

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Company	<b>Renu Medical, Inc.</b>
Location	Everett, WA, US
Contact	L BRUCE PIERSON
510(k) history	9 submissions · 9 cleared · 2003-2021

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)Device record: <https://www.510kdatabase.net/k111773/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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