

K181738 Medline ReNewal Reprocessed OxiMax Pulse Oximetry Sensors

Mar 22, 2019
263 days to decision

K181738 · Product code: **DQA** · Anesthesiology
Source: <https://www.510kdatabase.net/k181738/>

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Oximeter (DQA)
Date received	Jul 2, 2018
Decision date	Mar 22, 2019
Days to decision	263 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Surgical Instrument Service and Savings Inc (Dba Medline
Location	Redmond, OR, US
Contact	Stephanie Boyle Mays
510(k) history	1 submissions · 1 cleared · 2019-2019

510k Database - www.510kdatabase.net

Device record: <https://www.510kdatabase.net/k181738/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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