

K213933 Percent Oxygen SensorsAug 19, 2022
246 days to decisionK213933 · Product code: **CCL** · Anesthesiology
Source: <https://www.510kdatabase.net/k213933/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Analyzer, Gas, Oxygen, Gaseous-phase (CCL)
Date received	Dec 16, 2021
Decision date	Aug 19, 2022
Days to decision	246 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Careox, LLC
Location	Princeton, NJ, US
Contact	Young Kim
510(k) history	1 submissions · 1 cleared · 2022-2022

REGULATORY CONSULTANT

Consulting firm	ProMedic Consulting, LLC
Contact	Paul Dryden

Regulatory consulting firm that managed this 510(k) submission on behalf of the applicant. Source: [FDA accessdata.fda.gov](https://accessdata.fda.gov)

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