

K211056 Oxus Sieve BedsMay 31, 2023
782 days to decisionK211056 · Product code: **CAW** · Anesthesiology
Source: <https://www.510kdatabase.net/k211056/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Generator, Oxygen, Portable (CAW)
Date received	Apr 9, 2021
Decision date	May 31, 2023
Days to decision	782 days
Third-party review	No
Combination product	No
PCCP authorized	No
Summary / Statement	Summary

APPLICANT

Company	Oxus, Inc.
Location	Auburn Hills, MI, US
Contact	Ryan Lenarcic
510(k) history	2 submissions · 2 cleared · 2017-2023

REGULATORY CONSULTANT

Consulting firm	Oxus, Inc. % Promedic, 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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