

K101154 CENTROX, RELIANT, ULTROX, AS-D+, AS-E, AS-G, AS-J, AS-K AND AS-LMar 28, 2011
339 days to decisionK101154 · Product code: **CAW** · Anesthesiology
Source: <https://www.510kdatabase.net/k101154/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Generator, Oxygen, Portable (CAW)
Date received	Apr 23, 2010
Decision date	Mar 28, 2011
Days to decision	339 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Airsep Corp.
Location	Buffalo, NY, US
Contact	PETER WEISENBORN
510(k) history	18 submissions · 17 cleared · 1987-2011

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k101154/>; Data sourced from FDA 510(k) public records (accessdata.fda.gov).

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