

**K001430 RESPIRTECH PRO-O2C, MODEL 2200 AND
RESPIRTECH PRO, MODEL 2000**Jul 5, 2000
61 days to decisionK001430 · Product code: **BTL** · Anesthesiology
Source: <https://www.510kdatabase.net/k001430/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Ventilator, Emergency, Powered (resuscitator) (BTL)
Date received	May 5, 2000
Decision date	Jul 5, 2000
Days to decision	61 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Vortran Medical Technology 1, Inc.
Location	Mchenry, IL, US
Contact	GORDON A WONG
510(k) history	28 submissions · 28 cleared · 1984-2021

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k001430/> Data sourced from FDA 510(k) public records (accessdata.fda.gov).

510k Database is an independent platform not affiliated with the U.S. Food and Drug Administration. Always verify at accessdata.fda.gov. - Generated July 9, 2026