

K962879 SALTERLABS 8660 SERIES NEBULIZEROct 22, 1996
90 days to decisionK962879 · Product code: **CAF** · Anesthesiology
Source: <https://www.510kdatabase.net/k962879/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nebulizer (direct Patient Interface) (CAF)
Date received	Jul 24, 1996
Decision date	Oct 22, 1996
Days to decision	90 days
Third-party review	No
Summary / Statement	Summary

APPLICANT

Company	Salter Labs
Location	Mchenry, IL, US
Contact	JAMES N CURTI
510(k) history	48 submissions · 48 cleared · 1978-2017

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k962879/> 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 May 24, 2026