

K790154 MANIFOLD, FOR DOUBLE SUPPLYMar 7, 1979
54 days to decisionK790154 · Product code: **CAF** · Anesthesiology
Source: <https://www.510kdatabase.net/k790154/>**SUBMISSION DETAILS**

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nebulizer (direct Patient Interface) (CAF)
Date received	Jan 12, 1979
Decision date	Mar 7, 1979
Days to decision	54 days
Third-party review	No

APPLICANT

Company	Airlife, Inc.
Location	Mchenry, IL, US
510(k) history	76 submissions · 76 cleared · 1979-1984

510k Database - www.510kdatabase.netDevice record: <https://www.510kdatabase.net/k790154/> 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 27, 2026