FDA 510(k) Premarket Notification

K830895 SINGLE USE NEBULIZER, ADJUST. AIR-

CLEARED

Apr 28, 1983
38 days to decision

K830895 · Product code: **CAF** · Anesthesiology
Source: https://www.510kdatabase.net/k830895/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Nebulizer (direct Patient Interface) (CAF)
Date received	Mar 21, 1983
Decision date	Apr 28, 1983
Days to decision	38 days
Third-party review	No

APPLICANT

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