FDA 510(k) Premarket Notification

K790154 MANIFOLD, FOR DOUBLE SUPPLY

CLEARED

Mar 7, 1979
54 days to decision

K790154 · 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