FDA 510(k) Premarket Notification

K810947 AIR ENTRAINMENT TRACH T

CLEARED

Jun 2, 1981
56 days to decision

K810947 · Product code: **BYF** · Anesthesiology
Source: https://www.510kdatabase.net/k810947/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Mask, Oxygen, Low Concentration, Venturi (BYF)
Date received	Apr 7, 1981
Decision date	Jun 2, 1981
Days to decision	56 days
Third-party review	No

APPLICANT

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