FDA 510(k) Premarket Notification

K013251 SPIRIT 300

CLEARED

Dec 18, 2001
81 days to decision

K013251 · Product code: **BYJ** · Anesthesiology
Source: https://www.510kdatabase.net/k013251/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Unit, Liquid-oxygen, Portable (BYJ)
Date received	Sep 28, 2001
Decision date	Dec 18, 2001
Days to decision	81 days
Third-party review	Yes
Summary / Statement	Summary

APPLICANT

Company	**Caire, Inc.**
Location	Littleton, CO, US
Contact	ROGER BRIESE
510(k) history	7 submissions · 7 cleared · 1995-2025