FDA 510(k) Premarket Notification

K800546 BUNN VOLUMAIRE RESPIRATORY EXERCISER

CLEARED

Mar 25, 1980
15 days to decision

K800546 · Product code: **BWF** · Anesthesiology
Source: https://www.510kdatabase.net/k800546/

SUBMISSION DETAILS

Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spirometer, Therapeutic (incentive) (BWF)
Date received	Mar 10, 1980
Decision date	Mar 25, 1980
Days to decision	15 days
Third-party review	No

APPLICANT

Company	**The John Bunn Co.**
Location	Mchenry, IL, US
510(k) history	20 submissions · 20 cleared · 1976-1990