

**K800546 BUNN VOLUMAIRE RESPIRATORY EXERCISER**Mar 25, 1980  
15 days to decisionK800546 · Product code: **BWF** · Anesthesiology  
Source: <https://www.510kdatabase.net/k800546/>**SUBMISSION DETAILS**

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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**

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

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510k Database - [www.510kdatabase.net](http://www.510kdatabase.net)

Device record: <https://www.510kdatabase.net/k800546/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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