

**K833337 VOLUREX II**Nov 28, 1983  
62 days to decisionK833337 · Product code: **BWF** · AnesthesiologySource: <https://www.510kdatabase.net/k833337/>**SUBMISSION DETAILS**

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Decision	Substantially Equivalent (Cleared)
Submission type	Traditional
Device classification	Spirometer, Therapeutic (incentive) (BWF)
Date received	Sep 27, 1983
Decision date	Nov 28, 1983
Days to decision	62 days
Third-party review	No

**APPLICANT**

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Company	<b>Dhd Medical Products Div. Diemolding Corp.</b>
Location	Mchenry, IL, US
510(k) history	43 submissions · 43 cleared · 1979-1988

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

Device record: <https://www.510kdatabase.net/k833337/> Data sourced from FDA 510(k) public records ([accessdata.fda.gov](http://accessdata.fda.gov)).

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